

treatment; Neisseria infection; meningitis; septicaemia; gonorrhea.

XX OS Neisseria gonorrhoeae.
XX PN W09924578-A2.
XX PD 20-MAY-1999.
XX PF 09-OCT-1998; 98WO-IB01665.
XX PR 01-SEP-1998; 98GB-0019016.
XX PR 06-NOV-1997; 97GB-0023516.
XX PR 14-NOV-1997; 97GB-0024190.
XX PR 18-NOV-1997; 97GB-0024386.
XX PR 27-NOV-1997; 97GB-0025158.
XX PR 10-DEC-1997; 97GB-0026147.
XX PR 14-JAN-1998; 98GB-0000759.
XX PA (CHIR-) CHIRON SPA.
XX PI Grandi G, Masignani V, Pizza M, Rappuoli R, Scarlato V;
XX N-PSDB; AA212028.
XX DR WPI: 1999-327407/27.
XX DR N-PSDB; AA212028.
XX PT Proteins from Neisseria meningitidis and N. gonorrhoeae useful for
XX diagnosis, treatment and prevention of infection
XX PS Claim 4; Page 125; 524pp; English.
XX CC Amino acid sequences AAY38499-Y38944 represent Neisseria meningitidis
XX and N. gonorrhoeae antigenic proteins. They are encoded by open
XX reading frames (ORFs) AX211972-Z12358. The antigenic proteins,
XX their fragments, their nucleic acids and antibodies are used for
XX diagnosis, prevention (as vaccines) or treatment of Neisseria
XX infections, such as meningitis, septicaemia and gonorrhea. Both
XX organisms are closely related. Fragments of the nucleic acids
XX are useful as hybridisation probes and antisense reagents.
XX SX Sequence 447 AA;

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Ratio: 5.103 Gaps: 0
Percent Similarity: 100.000 Percent Identity: 100.000

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ID AAV38561 standard; Protein: 447 AA.

AC AAV38561;

DT 08-OCT-1999 (first entry)

Neisseria meningitidis antigen encoded by ORF22.

Neisseria meningitidis; Neisseria gonorrhoeae; antigen; vaccine;
 treatment; Neisseria infection; meningitis; septicaemia; gonorrhea.

Neisseria meningitidis.

W09924578-A2.

20-MAY-1999.

09-OCT-1998; 98WO-IB01665.

01-SEP-1998; 98GB-0019016.

06-NOV-1997; 97GB-0023516.

14-NOV-1997; 97GB-0024190.

18-NOV-1997; 97GB-0024386.

27-NOV-1997; 97GB-0025158.

10-DEC-1997; 97GB-0026147.

14-JAN-1998; 98GB-0000759.

(CHIR-) CHIRON SPA.

Grandi G, Masignani V, Pizza M, Rappuoli R, Scarlato V;

WPI; 1999-327407/27.

N-PSDB; AA212026.

Proteins from Neisseria meningitidis and N. gonorrhoeae useful for
 diagnosis, treatment and prevention of infection

Claim 4; Page 123; 524pp; English.

Amino acid sequences AAV38499-Y38944 represent Neisseria meningitidis
 and N. gonorrhoeae antigenic proteins. They are encoded by open
 reading frames (ORFs) AA211972-Z12358. The antigenic proteins,
 their fragments, their nucleic acids and antibodies are used for
 diagnosis, prevention (as vaccines) or treatment of Neisseria
 infections, such as meningitis, septicaemia and gonorrhea. Both
 organisms are closely related. Fragments of the nucleic acids
 are useful as hybridisation probes and antisense reagents.

Sequence 447 AA;

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 Ratio: 4.980 Gaps: 0
 Percent Similarity: 99.776 Percent Identity: 96.197

alignment_block:

US-09-303-518D-131 x AAY38561 ..

Align seg 1/1 to: AAY38561 from: 1 to: 447

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XX DT 08-OCT-1999 (first entry)
XX DE Neisseria meningitidis strain A antigen encoded by ORF22.
XX KW Neisseria meningitidis; Neisseria gonorrhoeae; antigen; vaccine;
XX KW treatment; Neisseria infection; meningitis; septicaemia; gonorrhea.
XX OS Neisseria meningitidis.
XX PN WO924578-A2.
XX PD 20-MAY-1999.
XX PF 09-OCT-1998; 98WO-1801665.
XX PR 01-SEP-1998; 98GB-0019016.
XX PR 06-NOV-1997; 97GB-0023516.
XX PR 14-NOV-1997; 97GB-0024190.
XX PR 18-NOV-1997; 97GB-0024386.
XX PR 27-NOV-1997; 97GB-0025158.
XX PR 10-DEC-1997; 97GB-0026147.
XX PR 14-JAN-1998; 98GB-0000759.
XX PA (CHIR-) CHIRON SPA.
XX Grandi G, Masignani V, Pizza M, Rappuoli R, Scarlato V;
PI
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XX WPI; 1999-327407/27.
DR N-PSDB; AA212027.
XX Proteins from Neisseria meningitidis and N. gonorrhoeae useful for
PT diagnosis, treatment and prevention of infection
PS Claim 4; Page 123; 524pp; English.
XX Amino acid sequences AAY38499-Y38944 represent Neisseria meningitidis
CC and N. gonorrhoeae antigenic proteins. They are encoded by open
CC reading frames (ORFs) AA211972-212358. The antigenic proteins,
CC their fragments, their nucleic acids and antibodies are used for
CC diagnosis, prevention (as vaccines) or treatment of Neisseria
CC infections, such as meningitis, septicaemia and gonorrhea. Both
CC organisms are closely related. Fragments of the nucleic acids
CC are useful as hybridisation probes and antisense reagents.
XX SQ Sequence 447 AA;
alignment_scores:
Quality: 2148.00 Length: 447
Ratio: 4.927 Gaps: 0
Percent Similarity: 97.539 Percent Identity: 93.289
alignment_block:
US-09-303-518D-131 x AAY38562 ..
Align seg 1/1 to: AAY38562 from: 1 to: 447
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ID AA1999 standard; Protein; 322 AA.

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DT 08-OCT-1999 (first entry)
XX Neisseria gonorrhoeae antigen encoded by a partial ORF22.
DE
XX Neisseria meningitidis; Neisseria gonorrhoeae; antigen; vaccine;
KW treatment; Neisseria infection; meningitis; septicaemia; gonorrhea.
XX
OS Neisseria gonorrhoeae.
XX
PN WO9924578-A2.
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PD 20-MAY-1999.
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PF 09-OCT-1998; 98WO-IB01665.
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PR 01-SEP-1998; 98GB-0019016.
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PR 14-NOV-1997; 97GB-0024190.
PR 18-NOV-1997; 97GB-0024386.
PR 27-NOV-1997; 97GB-0025158.
PR 10-DEC-1997; 97GB-0026147.
PR 14-JAN-1998; 98GB-0000759.
XX
PA (CHIR-) CHIRON SPA.
XX
PI Grandi G, Masignani V, Pizza M, Rappuoli R, Scarlato V;
XX WPI; 1999-327407/27.
XX
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Proteins from Neisseria meningitidis and N. gonorrhoeae useful for diagnosis, treatment and prevention of infection

Claim 4; Page 124-125; 524pp; English.

Amino acid sequences AAY38499-Y38944 represent Neisseria meningitidis and N. gonorrhoeae antigenic proteins. They are encoded by open reading frames (ORFs) AA11972-212358. The antigenic proteins, their fragments, their nucleic acids and antibodies are used for diagnosis, prevention (as vaccines) or treatment of Neisseria infections, such as meningitis, septicaemia and gonorrhea. Both organisms are closely related. Fragments of the nucleic acids are useful as hybridisation probes and antisense reagents.

Sequence 322 AA;

alignment_scores:

Quality: 1631.00 Length: 322

Ratio: 5.065 Gaps: 0

Percent Similarity: 100.000 Percent Identity: 100.000

alignment_block:

US-09-303-518D-131 x AAY38563 ..

Align seg 1/1 to: AAY38563 from: 1 to: 322

1 ATGATTAATAAATAAAGGTCTAAATCTGCCCATCGCGGACAGCGGA 50

1 MetileLysileLysLysGlyLeuAsnLeuProIleAlaGlyArgProG 17

51 GCAAGTCATTATGACGCGCGCGCATACCGAAGTCGCGTTCGTCGGCG 100

17 uGlnValIleTyrAspGlyProAlaIleThrGluValAlaLeuLeuGly 34

101 AAGAATATGTCGCGCATCGCCCTCGATGAAATCAAGAAAGTGAAGCC 150

34 lueLysrValGlyMetArgProSerMetLysileLysGlyGluAla 50

151 GTCAAAAAGGCGCAAGTGTCTGTTTGAAGACAAAAGATCCGCGCGTAGT 200

51 ValLysLysGlyGlnValLeuPheGluAspLysLysAsnProGlyValVa 67

201 ATTTACTGCGCGCGCTTCAGGCAAAATCCCGCTATTACCGTGGCGAA 250

251 AGCGCTACTTCACTCAGTCGTGATTGCGGTGAGCGCAACGACGAAATC 300
 |||||
 84 ysargValLeuGlnSerValValleAlaValGlu**AsnAspGluIle 100
 |||||
 301 GAGTTCGACGCTACTGACGCTGAGCGCTGGCAAAATTCAGCAGCGAAA 350
 |||||
 101 GluPheGluArgTyrAlaProGluAlaLeuAlaAsnLeuSerGlyGluG 117
 |||||
 351 AGTGGCCCGCAACCTGATCAATCAGGCTTATGGACTGCGCTTCGACCC 400
 |||||
 117 uValArgArgAsnLeuIleGlnSerGlyLeuTrpThrAlaLeuArgThra 134
 |||||
 401 GTCCGTTACGAAATCCCTGCGTAGATGCGGAGCGCTTCGCCATCTTC 450
 |||||
 134 rgProPheSerLysIleProAlaValAspAlaGluProPheAlaIlePhe 150
 |||||
 451 GTCATGCGGATGGACACCAATCCG 474
 |||||
 151 ValAsnAlaMetAspThrAsnPro 158

seq_name: /SIBS1/gcgdata/geneseq/gene-seq-embl/AA1999.DAT.AAY34439

seq_documentation_block:

ID AAY34439 standard; Protein; 451 AA.

AC AAY34439;

DT 25-AUG-1999 (first entry)

DE Porphyromonas gingivalis protein PGI.

KW Porphyromonas gingivalis; PG; periodontal disease; gingivitis;
 KW vaccine; antigenic.

OS Porphyromonas gingivalis.

PN WO9929870-A1.

PD 17-JUN-1999.

PF 10-DEC-1998; 98WO-AU01023.

PR 04-AUG-1998; 98AU-0005028.

PR 10-DEC-1997; 97AU-0000839.

PR 31-DEC-1987; 97AU-0001182.

PR 30-JAN-1998; 98AU-0001546.

PR 10-MAR-1998; 98AU-0002264.

PR 09-APR-1998; 98AU-0002911.

PR 23-APR-1998; 98AU-0003128.

PR 05-MAY-1998; 98AU-0003338.

PR 22-MAY-1998; 98AU-0003654.

PR 29-JUL-1998; 98AU-0004917.

XX (CSLC-) CSL LTD.

XX Agius CT, Barr IG, Hocking DM, Margetts MB, Patterson MA;

XX Ross BC, Rothel LJ, Webb EA;

XX WPI; 1999-385613/32.

XX N-PSDB; AAX91657.

XX Antigenic Porphyromonas gingivalis peptides for preventing

XX gingivitis

XX Claim 1; Page 417-418; 588pp; English.

XX AAX91536 to AAX91801 encode two hundred and sixty six antigenic

CC Porphyromonas gingivalis (PG) polypeptide sequences given in AAY34318 to

CC AAY34583, AAX91802 to AAX91989 represent PCR primers used in the

CC isolation of the PG polypeptides. The PG polypeptides have antibacterial

CC activity with a vaccine mechanism of action. The PG polypeptides can be

CC used as vaccines especially against Porphyromonas gingivalis. Probes can

CC be used to detect Porphyromonas gingivalis in standard hybridisation

CC assays. Porphyromonas gingivalis is involved in periodontal disease

CC especially gingivitis.

XX SQ Sequence 451 AA;

alignment_scores: Quality: 648.00 Length: 452

Ratio: 2.189 Gaps: 7

Percent Similarity: 65.487 Percent Identity: 34.071

alignment_block:

US-09-303-518d-131 x AAY34439 ..

Align seg 1/1 to: AAY34439 from: 1 to: 451

1 ATGATTAAATCAAAAAAGTCTAAATCTGCCCATCGCGGACACCGGA 50
 :|||
 4 ValIleLysThrLysLysGlyLeuAlaLeuAlaLeuLysGlyLysProLe 20
 :|||
 51 GCAAGTCATTTATGACGCGCCGCTTACCGAAGTC...GCGTTGCTTG 97
 :|||
 20 uProGluMetLeuAlaGluProAlaGlnSerProThrTyrAlaValValp 37
 :|||
 98 GCGAAGAATATGTCGCGCATGCGCCCTCGATGAAATCAAGGAGGTGAA 147
 :|||
 37 roAspAspPheGluGlyValIleProLysValThrAlaArgProGlyAsp 53
 :|||
 148 GCCGTCAAAAAAGCCCAAGTCTGTTTGAAGACAAAAAAGAAATCGGCGCT 197
 :|||
 54 LysValArgAlaGlySerAlaLeuMetHisLysAlaTyrProGluMe 70
 :|||
 198 AGATTATTTCGCGCGCTTCAGGCAAAATCGCGCTATTACCGTGGCG 247
 :|||
 70 tLysPheThrSerProValSerGlyGluValIleAlaValAsnArgGlyA 87
 :|||
 248 AAAAGCGCTACTTCAGTCAGTGTGCTGCTGCTGTAAGGCAACGACGAA 297
 :|||
 87 laLysArgLysValLeuSerIleGluValLysProAspGlyLeuAsnGlu 103
 :|||
 298 ATCGAG...TTCGACGCTACGTACCTGAGCGCTGGCAAAATTCGACGAG 344
 :|||
 104 TyrGluSerPheProValGlyAspProSerAla.....LeuSerAl 117
 :|||
 345 CGAAAAAGTCGCGCGCAACCTGATTCAATCAGCTTATGGACTGCGCTTC 394
 :|||
 117 aGluGlnIleLysGluLeuLeuSerSerGlyMetTrpGlyPheIleLe 134
 :|||
 395 GCACCCGCTCGTTTCAGCAAAATCCCTGCGGTAGATCGCGCGCTTCGCC 444
 :|||
 134 yGlnArgProTyrAspIleValAlaThrProAspIleAlaProArgAsp 150
 :|||
 445 ATCTTCGTCAATCGGATGACACCAATCCGCTGCGCGACCCCTACGCT 494
 :|||
 151 IleTyrIleThrAlaAsnPheThrAlaProLeuAlaProAspPheAspPh 167
 :|||
 495 CATCATCAAGAAGCGCGCAAGACTTCAACGCGCTGTTGGTATTCA 544
 :|||
 167 eIleValArgGlyGluGluArgAlaLeuGlnThrAlaIleAspAlaLeu 184
 :|||
 545 GCGCGCTGACCGCAACGTAATCCATGTGTGTAAGCAGCAGCGCGCAGAC 594
 :|||
 184 laLysLeuThrThrGlyLysValTyrValGlyLeuLysProGlySerSer 200
 :|||
 595 GTGCGCTCTGAAATGCTCCCAATATCCAAACACATGAATTTGGCGGCC 644
 :|||
 201 LeuGlyLeuHisAsnAlaGluIleValGluValHis.....GlyPr 214
 :|||
 645 GCATCTCTGCGCGCTTGTAGTGGCAGCAGCATTCATTTCATCGAGCAGTCG 694
 :|||
 214 ohisProAlaGlyAsnValGlyValIleAsnHisThrLysProIleA 231

```
695 GCGGCAATAAACCGTGTGGACCATCAATTATCAAGACGTGATTGCTATC 744
:: : : : : : : : : : : : : : : : : : : : : : : : : : : : : :
231 snArgGlyGluThrValTrpThrLeuLysAlaThrAspLeuIleValIle 247
:: : : : : : : : : : : : : : : : : : : : : : : : : : : : : :
745 GGAGCTTTGTTCTGTAACAGCGCTCTGAATACCGAGCGGTGGTGGCTT 794
:: : : : : : : : : : : : : : : : : : : : : : : : : : : : : :
248 GlyArgPheLeuLeuThrGlyLysAlaAspPheThrArgMetIleAlaMe 264
:: : : : : : : : : : : : : : : : : : : : : : : : : : : : : :
795 GGGGGCGCTGCAAGTCAACAAACCGCGCTCTTGGCTGACCTTTGGGTG 844
:: : : : : : : : : : : : : : : : : : : : : : : : : : : : : :
264 tThrGlySerAspAlaAlaHisGlyTyrValArgIleMetProGlyC 281
:: : : : : : : : : : : : : : : : : : : : : : : : : : : : : :
845 CGAAGGTGTCTCACTTACCGCGCGCAATTGGTT.....GACGCGGAC 888
:: : : : : : : : : : : : : : : : : : : : : : : : : : : : : :
281 yAsnValPheAlaSerPheProGlyArgLeuThrIleLysGluSerHis 297
:: : : : : : : : : : : : : : : : : : : : : : : : : : : : : :
889 AACCGCTGATTTCGGTTCGGTATTGAACGGTTCGATTGCACAAAGCGC 938
:: : : : : : : : : : : : : : : : : : : : : : : : : : : : : :
298 GluArgValIleAspGlyAsnValLeuThrGlyLysLysLeuGluLys 314
:: : : : : : : : : : : : : : : : : : : : : : : : : : : : : :
939 GCATGATTATTGGACGCTACCAACATCAGATTCCGTTATCGAAGAAG 988
:: : : : : : : : : : : : : : : : : : : : : : : : : : : : : :
314 sGluProPheLeuSerAlaArgCysAspGlnIleThrValIleProGlu 331
:: : : : : : : : : : : : : : : : : : : : : : : : : : : : : :
989 GCCGCGAGC...AAAGAGCTGTTCGGCTGGTTCGCGCGACGCGGACAAA 1035
:: : : : : : : : : : : : : : : : : : : : : : : : : : : : : :
331 lyAspAspValAspGluLeuPheGlyTrpAlaAlaProArgLeuAspGln 347
:: : : : : : : : : : : : : : : : : : : : : : : : : : : : : :
1036 TACTCCATCAGCGCACCTCTCGGCCATTTCCTA...AAACAACAACT 1082
:: : : : : : : : : : : : : : : : : : : : : : : : : : : : : :
348 TyrSerMetSerArgAlaTyrPheSerTrpLeuGlnGlyLysAsnLysG 364
:: : : : : : : : : : : : : : : : : : : : : : : : : : : : : :
1083 CTTCAAGTTCACGACCGCTCAACGCGCGACCGCGCGCTGATACCGGA 1132
:: : : : : : : : : : : : : : : : : : : : : : : : : : : : : :
364 uTyrValLeuAspAlaArgIleLysGlyGlyAlaArgAlaMetIleMetS 381
:: : : : : : : : : : : : : : : : : : : : : : : : : : : : : :
1133 TCGCCTATTAGCGCGGTAAATCGGTGGACATCTCGCTACCTTGCCTT 1182
:: : : : : : : : : : : : : : : : : : : : : : : : : : : : : :
381 eAsnGluTyrAspArgValPheProMetAspIleTyrProGluTyrLeu 397
:: : : : : : : : : : : : : : : : : : : : : : : : : : : : : :
1183 TTGCGCGATTAAATCGTGGCGATACGACGCGCGAGGCTTGGGTG 1232
:: : : : : : : : : : : : : : : : : : : : : : : : : : : : : :
398 LeuLysAlaIleIleAlaPheAspIleAspLysMetGluAspLeuGlyI 414
:: : : : : : : : : : : : : : : : : : : : : : : : : : : : : :
1233 CTTGGAATTGGACGAAGACCTCGCTTGTGACGCTTCGCTGCGCGG 1282
:: : : : : : : : : : : : : : : : : : : : : : : : : : : : : :
414 eTyrGluValAlaProGluAspPheAlaThrCysGluPheValAspThrS 431
:: : : : : : : : : : : : : : : : : : : : : : : : : : : : : :
1283 GCATATAGATACGCGCGCTGTGCGCAAGTCTGCGGAACCATTCAGT 1332
:: : : : : : : : : : : : : : : : : : : : : : : : : : : : : :
431 eRlySileGluLeuGlnArgIleValArgGluGlyLeuAspMetLeuTyr 447
:: : : : : : : : : : : : : : : : : : : : : : : : : : : : : :
1333 AAGGAA 1338
:: : : : : : : : : : : : : : : : : : : : : : : : : : : : : :
448 LysGlu 449
```

seq_name: /SIDSL/gcdata/geneseq/geneseq-emb1/AA1999.DAT:AAAY34318

seq_documentation_block:

ID AAAY34318 standard; Protein; 454 AA.

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XX 17-JUN-1999.
PD 10-DEC-1998; 98WO-AU01023.
XX 04-AUG-1998; 98AU-0005028.
PR 10-DEC-1997; 97AU-0000839.
PR 31-DEC-1997; 97AU-0001182.
PR 30-JAN-1998; 98AU-0001546.
PR 10-MAR-1998; 98AU-0002264.
PR 09-APR-1998; 98AU-0002911.
PR 23-APR-1998; 98AU-0003128.
PR 05-MAY-1998; 98AU-0003338.
PR 22-MAY-1998; 98AU-0003654.
PR 29-JUL-1998; 98AU-0004917.
XX (CSLC-) CSL LTD.
FA Agius CT, Barr IG, Hocking DM, Margetts MB, Patterson MA;
PI Ross BC, Rothel LJ, Webb EA;
XX WPI: 1999-385613/32.
DR N-PSDB; AAX91536.
XX Antigenic Porphyromonas gingivalis peptides for preventing
PT gingivitis
XX Claim 1; Page 277; 588pp; English.
XX AAX91536 to AAX91801 encode two hundred and sixty six antigenic
CC Porphyromonas gingivalis (PG) polypeptide sequences given in AAY34318 to
CC AAY34583. AAX91802 to AAX91989 represent PCR primers used in the
CC isolation of the PG polypeptides. The PG polypeptides have antibacterial
CC activity with a vaccine mechanism of action. The PG polypeptides can be
CC used as vaccines especially against Porphyromonas gingivalis. Probes can
CC be used to detect Porphyromonas gingivalis in standard hybridisation
CC assays. Porphyromonas gingivalis is involved in periodontal disease
CC especially gingivitis.
XX Sequence 454 AA;
```

alignment_scores:

Quality: 648.00 Length: 452

Ratio: 2.189 Gaps: 7

Percent Similarity: 65.487 Percent Identity: 34.071

alignment_block:

US-09-303-518D-131 x AAY34318 ..

Align seg 1/1 to: AAY34318 from: 1 to: 454

1 ATGATTAAATCAAAAAGGTCTAAATCTGCCATCGCGGCGACGCCGA 50

:::||||| ||||| ||| ::|::|::|

7 ValIleLysThrLysLysGlyLeuAlaLeuAsnLeuLysGlyLysProLe 23

:::|::|::| ||||| ||| ::|::|::|

51 GCAAGTCATTATGACGCGCGCCCATACCGAAGTC...GCGTTCCTTG 97

::: ||||| ::|::|::|

23 uProGluMetLeuAlaGluProAlaGlnSerProThrTyrAlaValP 40

98 GCGAAGAATATGTCGCATCGCCCTCGATGAAATCAAGGAAGTGAA 147

:::|::|::| ||||| ||| ::|::|::|

40 roAspAspPheGluGlyValIleProLysValThrAlaArgProGlyasp 56

:::|::|::| ||||| ||| ::|::|::|

148 GCCGTCAAAAGGCCAAGTGTGTTGAAGACAAAGAAATCCGCGCT 197

||::|::| |||||::|::|::| ||| ||| ::|::|::|

57 LysValArgAlaGlySerAlaLeuMetHisHisLysAlaTyrProGluMe 73

198 AGTATTACTGCGCGCTTCAGGCAAAATCGCGCTATTTCACCGTGGC 247

:|||||::|::|::|::|::|::|::|::|::|::|::|::|::|::|::|::|

73 tLysPheThrSerProValSerGlyGluValIleAlaValAsnArgGlyA 90

::|::|::|::|::|::|::|::|::|::|::|::|::|::|::|::|

248 AAAAGCGCGTACTTCAGTCAGTCGCTGATTGCCGTTGAAGCAACGACGAA 297

90 laYsArgLYsValLeuSerIleGluValLysProAspGlyLeuAsnGlu 106
 298 ATCGAG...TTCGAACGCTACGTACCTTGAAGCGCTGGCAAAATTTAGCAG 344
 107 TyrGluSerPheProValGlyAspProSerAla.....LeuSerAl 120
 345 CGAAAAAGTCGGCGCAACCTGATTCAATCAGCTTATGGACTGGCGTTC 394
 120 aGluInIleLysGluLeuLeuSerSerGlyMetTrpGlyPheIleL 137
 395 GCACCGCTCGGTTCCAGAAAATCCCTGCCGTAGATCCCGAGCGGTTCGC 444
 137 ysGlnArgProTyrAspIleValAlaThrProAspIleAlaProArgasp 153
 445 ATCTTCGTCATCGCATGGACCAATCCGCTGGCTGCGCGCACCCACCGT 494
 154 IleTyrIleThrAlaAsnPheThrAlaProLeuAlaProAspPheAspPh 170
 495 CATCATCAAAAGAGCCCGCAAGACTTCAACGGCGCTGTGGTATTGCA 544
 170 eIleValArgLYcLugluArgAlaLeuGlnThrAlaIleAspAlaLeuA 187
 545 GCCCGCTGACCGCAACGTAAATCCATCGTGTAAAGCAGCAGCGCGCAG 594
 187 laYsLeuThrThrGlyLysValTyrValcLYLeuLysProGlySerSer 203
 595 GTGCGGTCTCAAAATGCTCCAATATCGAAACACATGAATTTGGCGGCC 644
 204 LeuGlyLeuHisAsnAlaGluIleValGluValHis.....GlyPr 217
 645 GCATCTCGCGGCTGAGTGGCAGCAGCACATTCATTCATCGAGCGACGTCG 694
 217 oHisProAlaGlyAsnValcLYValLeuIleAsnHisThrLysProIleA 234
 695 GCGCGAATAAACCGTCTGGACCATCAATTATCAAGACGTGATGCTATC 744
 234 snArgGlyGluThrValTrpThrLeuLysAlaThrAspLeuIleValIle 250
 745 GGACGTTGTGCTGAACGCGCTCTGAATACCGCGCTGCTGCTACCGTTTGGGT 794
 251 GlyArgPheLeuLeuThrGlyLysAlaAspPheThrArgMetIleAlaMe 267
 795 GGCGGCGCTGCAAGTCAACAACCGCGCTCTGCTGCTACCGTTTGGGT 844
 267 tThrGlySerAspAlaAlaHisGlyTyrValArgIleMetProGlyC 284
 845 CGAAGTGTCTCAACTTACCGCGCGCAATTTGGT.....GAGCGCGAC 888
 284 ysAsnValPheAlaSerPheProGlyArgLeuThrIleLysGluSerHis 300
 889 AACCGGTGATTTCCGTTTCGGTATTGAACGTCGCGATTCACAAAGCGC 938
 301 GluArgValIleAspGlyAsnValLeuThrGlyLysLysLeuCysGluLY 317
 939 GCATGATTTATTTGGACGCTACCACAAATCAGATTTCGTTATTCGAAGA 988
 317 sGluProPheLeuSerAlaArgCysAspGlnIleThrValIleProGlu 334
 989 GCGCGCAGC...AAAGACTGTTTCGGCTGGGTGGCGCGCAGCCGACAAA 1035
 334 lYsAspAspValAspGluLeuPheGlyTrpAlaAlaProArgLeuAspGln 350
 1036 TACTCCATCAGCGCACCACTCTCGGCCATTTCCTA...AAACAACAAT 1082
 367 utYrValLeuAspAlaArgIleLysGlyGlyGluArgAlaMetIleMets 384
 1133 TCGGCACCTATCAGCGCGTATGCGTTGGACATCTCGCTACCTGCTT 1182

```

384 erAsnGluTy rAspArgValPheProMetAspIleTy rProGluTyLeu 400
1183 TTCCGCCGAATTAATCGTCGGCATACCGACAGCGCAGGCTTTGGGTG 1232
||||| :||::|| | ||::: |::| |::| |::| |::|
401 LeuLysAlaIlellealApeaspileaspIysMetGluaspleuGlyII 417
1233 CTTGGAAATTGGAGCAAGAAGACTCGCTTGTGCAGCTTGCTGCCCCG 1282
||||| |::| |::| |::| |::| |::| |::| |::|
417 eTy rGluValAlaIlaProGluAspPheAlaThrCysGluPheValAspThrs 434
1283 GCAAATACCAATACGCGCGCTGTGTGGCGCAAGAGTCTGGAACCATTCAG 1332
:||:| |::| |::| |::| |::| |::| |::| |::|
434 e rLy sIle dle uLeuGlnArgIleValArgGluGlyLeuAspMetLeuTy r 450
1333 AAGGAA 1338
|||||
451 LysGlu 452

seq_name: ./SIDS1/gcgdata/geneseq/geneseq-emb1/AA2000.DAT:AA Y75271
seq_documentation_block:
ID AA Y75271 standard; Protein; 119 AA.
XX AC
XX AC
XX AC
XX AC
XX 21-MAR-2000 (first entry)
XX DE Neisseria gonorrhoeae ORF 628 protein sequence SEQ ID NO:2016.
XX KW Neisseria meningitidis; Neisseria gonorrhoeae; antigen; vaccine;
KW K antigenic; diagnosis; immunogenic; infection; meningitis; septicaemia;
KW K antibacterial; gene therapy.
XX OS Neisseria gonorrhoeae.
XX XX
XX PN W09957280-A2.
XX PD
XX PF 11-NOV-1999.
XX PF 30-APR-1999; 99WO-US09346.
XX PF 01-MAY-1998; 98US-0083758.
XX PR 31-JUL-1998; 98US-0094869.
XX PR 02-SEP-1998; 98US-0098994.
XX PR 02-SEP-1998; 98US-0099062.
XX PR 09-OCT-1998; 98US-0103749.
XX PR 09-OCT-1998; 98US-0103794.
XX PR 09-OCT-1998; 98US-0103796.
XX PR 25-FEB-1999; 99US-0121528.
XX PA (CHIR ) CHIRON CORP.
XX PA (GENO-) INST GENOMIC RES.
XX PI Fraser C, Galeotti C, Grandi G, Hickey E, Masignani V, Mora M;
PI Petersen J, Pizza M, Rappuoli R, Ratti G, Scalato E, Scarselli M;
PI Tettelin H, Venter JC;
XX XX
XX WP1; 2000-062150/05.
XX N-PSDB; AA254033.
XX DR
XX PT Novel Neisserial polypeptides predicted to be useful antigens for
XX PS vaccines and diagnostics -
XX PS Claim 2; Page 1003; 1453pp; English.
XX CC
XX AA Z53015 to AA Z54536, AA Z54577 to AA Z54615, and AA Y74253 to AA Y75941
XX CC represent novel Neisseria meningitis and N. gonorrhoeae polynucleotides
XX CC and polypeptides AA Z54537 to AA Z54576 and AA Z54616 to AA Z5473 represe
XX CC PCR primers used in the exemplification of the present invention. The
XX CC polypeptides, the polynucleotides, antibodies and compositions of
XX CC the invention can be used as vaccines, as diagnostic reagents, and as
XX CC immunogenic compositions. The polypeptides can be used in the
XX CC manufacture of medicaments for treating or preventing infection due to

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CC Neisserial bacteria (e.g. meningitis and septicaemia), to detect the
CC presence of Neisseria bacteria, or to raise antibodies. They may also
CC be used to screen for agonists or antagonists, which may themselves
CC have use as antibacterial agents. The polynucleotides of the invention
CC may also be used in gene therapy protocols.

XX Sequence 119 AA;

alignment_scores:
Quality: 582.50 Length: 120
Ratio: 4.895 Gaps: 1
Percent Similarity: 99.167 Percent Identity: 99.167

alignment_block:

US-09-303-518D-131/rev x AAY75271

Align seg 1/1 to: AAY75271 from: 1 to: 119

674 ATGTGGTGGCCACTCAAGCGCGGAGGATCGGGCGCCAAATTCATGTGT 625
1 MetCysValProLeuLysProAlaGlyCysGlyProProAsnSerCysVa 17
624 TTCATATTTGCAGCATTTTCAGACGGCAGCTCTGCGCCTGCTGTAC 575
17 lSerIleLeuAlaAlaPheSerAspGlyThrSerAlaProAlaAlaLeuH 34
574 ACATGATGATTTACGTTTCGTCGAGCGGCTCAATACACAGCCCGGT 525
34 lThrTrpIleLeuArgSerValArgArgLeuAsnThrAsnArgProArg 50
524 TTGAAGTCTTCGGCGGCTTCTTCATGATGACCGTAGGTCGGCAGCAG 475
51 LeuLysSerSerAlaAlaSerLeuMetMetThrValGlySerAlaAlaSe 67
474 CGGATTTGTGTCCATTCGCAATTCAGCAAGATGGCAACGGCTCGCATCTA 425
67 rGlyLeuValSerIleAlaLeuThrLysMetAlaAsnGlySerAlaSerT 84
424 CGGAGGATTTTCTGTAACGGCGGGTGGCAAGCGAGTCCATTAAGCCT 375
84 hrAlaGlyIleLeuLeuAsnGlyArgValArgSerAlaValHisLysPro 100
374 GATTGAATCAGGTTTCGGCGCAGCTTTTTCGCTGCTCAATTTGCGCAGCG 325
101 Asp...lIeArgLeuArgArgThrPheSerLeuLeuAsnPheAlaSerAl 116
324 TTCAGGTACG 315
116 aSerGlyThr 119

seq_name: /SIDS1/gcgdata/geneseq/geneseq-emb1/AA2000.DAT:AA75272

seq_documentation_block:

ID AAY75272 standard; Protein; 120 AA.

XX AAY75272;

XX 21-MAR-2000 (first entry)

XX Neisseria meningitidis ORF 628 protein sequence SEQ ID NO:2018.

XX Neisseria meningitidis; Neisseria gonorrhoeae; antigen; vaccine;
KW antigenic; diagnosis; immunogenic; infection; meningitis; septicaemia;
KW antibacterial; gene therapy.

OS Neisseria meningitidis.

XX WO9957280-A2.

PN 11-NOV-1999.

XX 30-APR-1999; 99WO-US09346.

XX 01-MAY-1998; 98US-0083758.
PR 31-JUL-1998; 98US-0094869.
PR 02-SEP-1998; 98US-0098994.
PR 02-SEP-1998; 98US-0099062.
PR 09-OCT-1998; 98US-0103749.
PR 09-OCT-1998; 98US-0103794.
PR 09-OCT-1998; 98US-0103796.
PR 25-FEB-1999; 99US-0121528.

XX (CHIR) CHIRON CORP.
PA (GENO-) INST GENOMIC RES.

XX Fraser C, Galeotti C, Grandi G, Hickey E, Masignani V, Mora M;
PI Petersen J, Pizza M, Rappuoli R, Ratti G, Scalato E, Scarselli M;
FI Tettelin H, Venter JC;

XX WPI; 2000-062150/05.

DR N-PSDB; AA254034.

XX Novel Neisserial polypeptides predicted to be useful antigens for
PT vaccines and diagnostics

XX Claim 2; Page 1003; 1453pp; English.

XX AA253015 to AA254536, AA254577 to AA254615, and AAY74253 to AAY75941
CC represent novel Neisseria meningitis and N. gonorrhoeae polynucleotides
CC and polypeptides. AA254537 to AA254576 and AA254616 to AA255473 represent
CC PCR primers used in the exemplification of the present invention. The
CC polypeptides, the polynucleotides, antibodies and compositions of
CC the invention can be used as vaccines, as diagnostic reagents, and as
CC immunogenic compositions. The polypeptides can be used in the
CC manufacture of medicaments for treating or preventing infection due to
CC Neisserial bacteria (e.g. meningitis and septicaemia), to detect the
CC presence of Neisseria bacteria, or to raise antibodies. They may also
CC be used to screen for agonists or antagonists, which may themselves
CC have use as antibacterial agents. The polynucleotides of the invention
CC may also be used in gene therapy protocols.

XX Sequence 120 AA;

alignment_scores:

Quality: 545.00 Length: 119
Ratio: 4.698 Gaps: 0
Percent Similarity: 97.479 Percent Identity: 93.277

alignment_block:

US-09-303-518D-131/rev x AAY75272

Align seg 1/1 to: AAY75272 from: 1 to: 120

674 ATGTGGTGGCCACTCAAGCGCGGAGGATCGGGCGCCAAATTCATGTGT 625

1 MetCysValProLeuLysProAlaGlyCysGlyProProAsnSerCysVa 17

624 TTCATATTTGCAGCATTTTCAGACGGCAGCTCTGCGCCTGCTGTAC 575

17 lSerMetLeuAlaAlaPheSerAspGlyThrSerAlaProAlaAlaLeuG 34

574 ACATGATGATTTACGTTTCGTCGAGCGGCTCAATACACAGCGCGCGT 525

34 lThrTrpIleLeuArgSerValLysArgLeuAsnThrAsnArgProArg 50

524 TTGAAGTCTTCGGCGGCTTCTTCATGATGACCGTAGGTCGGCAGCAG 475

51 LeuLysSerSerAlaAlaSerLeuMetMetThrValGlySerAlaAlaSe 67

474 CGGATTTGTGTCCATTCGCAATTCAGCAAGATGGCAACGGCTCGCATCTA 425

67 rGlyLeuValSerIleAlaLeuThrLysMetAlaAsnGlySerAlaSerT 84

424 CGGAGGATTTTCTGTAACGGCGGGTGGCAAGCGAGTCCATTAAGCCT 375

|||||
84 hrAlaGlyIleLeuLeuAsnGlyArgValArgSerAlaValHisLysPro 100
374 GATTGAATCAGGTTGGCGGCACCTTTTCGCTGCCTCAATTTTCCAGCGC 325
||| ||||| ||||| ||| ||||| ||||| ||||| ||||| ||||| |||||
101 AspTrpIleLeuArgLeuArgThrSerProLeuLysPheAlaSerAl 117
324 TTCAGGT 318
|||||
117 aSerGly 119

seq_name: /SIDS1/gcgdata/geneseq/geneseq-emb1/AA2000.DAT:AAV75273

seq_documentation_block:

ID AAV75273 standard; Protein; 120 AA.

XX AC AAV75273;

XX DT 21-MAR-2000 (first entry)

XX DE Neisseria meningitidis ORF 628 protein sequence SEQ ID NO:2020.

XX KW Neisseria meningitidis; Neisseria gonorrhoeae; antigen; vaccine;
KW antigenic; diagnosis; immunogenic; infection; meningitis; septicaemia;
KW antibacterial; gene therapy.

XX OS Neisseria meningitidis.

XX PN W09957280-A2.

XX PD 11-NOV-1999.

XX PF 30-APR-1999; 99WO-US09346.

XX PR 01-MAY-1998; 98US-0083758.

XX PR 31-JUL-1998; 98US-0094869.

XX PR 02-SEP-1998; 98US-0098994.

XX PR 02-SEP-1998; 98US-0099062.

XX PR 09-OCT-1998; 98US-0103749.

XX PR 09-OCT-1998; 98US-0103794.

XX PR 09-OCT-1998; 98US-0103796.

XX PR 25-FEB-1999; 99US-0121528.

XX PA (CHIR) CHIRON CORP.

XX PA (GENO-) INST GENOMIC RES.

XX PI Fraser C, Galeotti C, Grandi G, Hickey E, Masignani V, Mora M;

XX PI Petersen J, Pizza M, Rappuoli R, Ratti G, Scalato E, Scarselli M;

XX PI Tettelin H, Venter JC;

XX DR WPI; 2000-062150/05.

XX DR N-PSDB; AA254035.

XX PT Novel Neisserial polypeptides predicted to be useful antigens for

XX PS vaccines and diagnostics

XX PS Claim 2; Page 1004; 1453pp; English.

XX CC AA253015 to AA254536, AA254577 to AA254615, and AA274253 to AAV75941
CC represent novel Neisseria meningitidis and N. gonorrhoeae polynucleotides
CC and polypeptides. AA254537 to AA254576 and AA254616 to AA254573 represent
CC PCR primers used in the exemplification of the present invention. The
CC polypeptides, the polynucleotides, antibodies and compositions of
CC the invention can be used as vaccines, as diagnostic reagents, and as
CC immunogenic compositions. The polypeptides can be used in the
CC manufacture of medicaments for treating or preventing infection due to
CC Neisserial bacteria (e.g. meningitis and septicaemia), to detect the
CC presence of Neisseria bacteria, or to raise antibodies. They may also
CC be used to screen for agonists or antagonists, which may themselves
CC have use as antibacterial agents. The polynucleotides of the invention
CC may also be used in gene therapy protocols.

XX SQ Sequence 120 AA;

alignment_scores:

Quality: 532.00 Length: 119

Ratio: 4.626 Gaps: 0

Percent Similarity: 96.639 Percent Identity: 89.916

alignment_block:

US-09-303-518D-131/rev x AAV75273 ..

Align seg 1/1 to: AAV75273 from: 1 to: 120

674 ATGTGCGTGCCTCAAGCGGCGAGGATGGCGGCCCAATTCATGTGT 625

|||||
1 MetCysValProLeuLysProAlaGlyCysGlyProProAsnSerCysVa 17

624 TTCGATATTTGGCAGCATTTTTCAGACGGCAGCTGTGGCGCTGCTGTAC 575

|||||
17 lSerMetLeuAlaAlaPheSeraspGlyThrSerAlaProAlaAlaLeuH 34

574 ACATCATGATTTTACGTTTCGCTCAGCGGCTCAATACCAACAGCGCGGT 525

|||||
34 lThrTrpIleLeuArgSerValLysArgLeuAsnThrSerLysProArg 50

524 TTGAAGTCTTCGCGGCTTCTTTTCATCATGACCGTAGGTCGGCAGCCAG 475

|||||
51 LeuLysSerSerAlaAlaSerLeuIleThrThrGlySerAlaAlaSe 67

474 CGGATTTGTTCCATCGCATTTGACGAAGATGGCGAAGCGGCTCGGCATCTA 425

|||||
67 rGlyLeuValSerIleAlaLeuThrLysMetAlaAsnGlySerAlaSerT 84

424 CGCAGGAGTTTTCGTCAGCAGCGGTCGGAAGCGCAGTCCTCAATAGCCT 375

|||||
84 hrAlaGlyIleLeuLeuAsnGlyArgValArgSerAlaValHisLysPro 100

374 GATTGAATCAGGTTGGCGGCACCTTTTTCGCTGCTCAATTTTCCAGCGC 325

|||||
101 AspTrpIleLeuArgLeuArgThrSerSerProLeuLysPheAlaAsnAl 117

324 TTCAGGT 318

|||||
117 aSerGly 119

seq_name: /SIDS1/gcgdata/geneseq/geneseq-emb1/AA2000.DAT:AAV82082

seq_documentation_block:

ID AAV82082 standard; Protein; 467 AA.

XX AC AAV82082;

XX DT 01-JUN-2000 (first entry)

XX DE Chlamydia pneumoniae antigen CPN100605 protein SEQ ID NO:2.

XX KW Chlamydia pneumoniae; antigen; CPN100605 protein; immunisation;

XX KW vaccine; infection; antibacterial; antiinflammatory; bronchitis;

XX KW community acquired pneumonia; upper respiratory tract infection;

XX KW sinusitis.

XX OS Chlamydia pneumoniae.

XX PN WO200006742-A2.

XX PD 10-FEB-2000.

XX PF 27-JUL-1999; 99WO-IB01331.

XX PR 27-JUL-1998; 98US-0094195.

XX PR 26-JUL-1999; 99US-0361443.

XX PA (CONN-) CONNAUGHT LAB LTD.

PI Murdin AD, Oomen RP;
 XX WPI; 2000-205466/18.
 DR N-PSDB; AAZ95378.
 XX Chlamydia pneumoniae antigens used for immunization and protection
 PT against Chlamydia diseases
 XX Claim 6; Fig 1; 48pp; English.

The present sequence represents the Chlamydia pneumoniae antigen
 CC CPN100605 protein. The CPN100605 protein has antibacterial and
 CC antiinflammatory activities. The Chlamydia pneumoniae CPN100605
 CC polynucleotide and protein can be used in vaccination methods for
 CC preventing and treating Chlamydia infection (e.g. infections caused by
 CC C. trachomatis, C. psittaci, C. pneumoniae or C. pecorum). The
 CC polynucleotide can be used to produce the protein recombinantly, in the
 CC construction of vaccine vectors, as a vaccine agent, and in the
 CC construction of an attenuated Chlamydia strain. The protein are also be
 CC useful as a vaccine agent, and for the preparation of medicaments for
 CC treating or preventing Chlamydia infection, e.g. community acquired
 CC pneumonia, and upper respiratory tract infections such as bronchitis and
 CC sinusitis.

XX Sequence 467 AA;

alignment_scores:
 Quality: 450.50 Length: 470
 Ratio: 1.532 Gaps: 16
 Percent Similarity: 62.553 Percent Identity: 29.149

alignment_block:

US-09-303-518D-131 x AAY82082 ..

Align seg 1/1 to: AAY82082 from: 1 to: 467

4 ATTAATAATCAAAAAGGCTAAATCTGCCATCGCGGACAGCG...GA 50
 3 IleThrValAsnArgGlyLeuAspLeuSerLeuGlnGlySerProLysG 19
 51 GCAAGTCATTTATGACGCGCGCGGCGCATACCGAAGTCGCGTGTTCGGC 100
 19 userGlyPheTyrAsn.....LysIleAsp 28
 101 AAGATATATGTCGC.....ATGCGCCG
 28 roGluPheValSerIleAspLeuArgProPheGlnProLeuSerLeuLys 44
 127 ATGAATAATCAAGGAGGTGAAGCGCTCAAAAAGGCCAAGTCGTGTTGA 176
 45 LeuLysValGluGlnGlyAspAlaValCysSerGlyAlaProIleAlaG 61
 177 AGACAAAAGAAATCGGCGGTAGTATTCTAGCGCGCGGCTTCAGGCAAA 226
 61 uTyrLysHisPheProAsnThrTyrIleThrSerHisValSerGlyVal 78
 227 TCGCGGTATTCACGCTGGCGAAAGCGGTACTTCAGTCAGTCGTGATT 276
 78 alThrAlaIleArgArgGlyAsnLysArgSerLeuLeuAspValIlelle 94
 277 ...GCCGTTGACGACGAGAAATCGAGTTCGAAGCTACGTACCTGA 323
 95 LysLysThrProGlyProThrSerThrGluTyr...ThrTyrAspLeuG 110
 324 AGCGTGGCAAAATGACACGCGAAAGTGCAGCGCGGCAACCTGATTCAAT 373
 110 nThrLeuSerArgSerAspLeuSerGluIlePheLys.....GluA 124
 374 CAGGCTTATGACTCGCTTCGACCGCTCGCTTCAGCAAAATCCCTGCC 423
 124 snGlyLeuPheAlaLeuIleLysGlnArgProPheAsp....IleProAla 139

424 GTAGATGCCGAG...CGTTGCCCATCTTCGTCAATCGCATGGACACAA 470
 140 ileProThrGlnThrProArgAspValPheIleAsnLeuAlaAspAsnAr 156
 471 TCCGCTGCTGCCACACCTAGGTATCATC.....A 502
 156 gProPheThrProSerProGluLysHisLeuAlaLeuPheSerSerArg 173
 503 AAGAACGCCGCGAAGACTTCAACGCCGCTGTTGGTATTGAGCGCGCTG 552
 173 luGluGlyPheTyrValPheValValGlyValArgAlaIleAlaLysLeu 189
 553 ACCGAACGTAATTCATGTGTAAAGCAGCAGCGCGCAGCGTCCGCTC 602
 190 PheGlyLeuAlaGProHisIleValPheArgAspArgLeuThrLeuProTh 206
 603 TCAAAATGCTGCCATATC...GAACACATGATGTTGGCGCGCGCATC 649
 206 rGlnGluLeuLysThrIleAlaHisLeuHisThrValSerGlyProPhe 223
 650 CTGCGCGCTTGAAGTGGCAGCAGCATTTTCATCGACGCGCGCGCG 699
 223 roSerGlySerProSerIleHisIleHisSerValAlaProIleThrAsn 239
 700 AATAAA...ACGCTGTGGACCATCAATATCAAGACGTGATGCTATCGG 746
 240 GluLysGluValValPheThrLeuSerPheGlnAspValLeuThrIleGl 256
 747 ACGTTTGTTCGTAACAGCGCGCTCGAATACCGAGCGCGTGGTTCCTGG 796
 256 yHisLeuPheLeuLysGlyArgIleLeuHisGluGlnValThrAlaLeuA 273
 797 GCGGC.....CTGCAAGTCAAAACCGCGCTCTTCGCGTACCGTTTGG 840
 273 laglyThrAlaLeuLysSerSerLeuArgArgTyrValIleThrLys 289
 841 GGTGGAGGTGCTCAACTTACCGCGCGGGAATGTTGAGCGGACAA 890
 290 GlyAlaSerPheSerSerLeuIleAsnLeuAsnAspIleSerAspAsnAs 306
 891 CGCGGTGATTTCGGTTCGTTGATGAACGGTTCGATTGCACAAAGCGCGC 940
 306 pThrLeuLysSerGlyAspProLeuThrGlyArgLeuLysLysLysGlu 323
 941 ATGAT...TATTGGGACGCTACCAATCAGATTTCGTTATCGAAGAA 987
 323 luGluProPheLeuGlyPheArgAspHisSerIleSerValLeuHisAsn 339
 988 GCGCGCAGCAAGAGCTGTCGCTGGTGGTGGCGCGCAGCGCGCAATA 1037
 340 ProThrLysArgGluLeuPheSerPheLeuArgIleGlyPheAsnLysPr 356
 1038 CTCATCAGCGCACCTCTCGCGCATTTCTCTAAATAACAACTCTTCA 1087
 356 ofrPheThrLysThrTyrLeuSerGlyPhePheLysLysLysArg...T 372
 1088 AGTTCAG.....ACAGCGCTCAACGCGCGCGCGCGCGCGCATGTA 1128
 372 hrTyrThrAsnProAspThrAsnLeuHisGlyGluThrArgProIlelle 388
 1129 CCGATCGGCATTTATGAGCGGTAAATCGCTGGTGGACATCTCGCTACCT 1178
 389 AspThrAspIleTyrAspLysValMetProMetArgIleProValValPr 405
 1179 GCTTTTGGCGATTTAATCGTCGGCGATACCGACAGCGCGCGCTTTGG 1228
 405 oLeuLysAlaValIleThrLysAsnPheAspLeuAlaAsnGluLeuG 422
 1229 GTTCTTGAATTTGACGAGAACCTCGCTTGTGACGCTTCGCTGCC 1278
 422 lypheLeuGluValCysGlyGluAspPheAlaLeuProThrLeuIleAsp 438
 1279 CCGCGCAATACGAATACGCGCGCTGTTGCGCAAAAGTCTGGAACCAT 1328

439 ProSerLysThrGluMetLeuThrIleValLysGluSerLeuLeuGluTyr 455
 1329 TGAGAGGAA 1338
 455 rAlaLysGlu 458

seq_name: /SIDS1/gcgdata/geneseq/geneseq-emb1/AA1999.DAT:AAV35375

seq_documentation_block:

ID AAV35375 standard; Protein; 469 AA.

XX AAV35375;

XX DT 13-SEP-1999 (first entry)

XX Chlamydia pneumoniae transmembrane protein sequence.

XX Respiratory disease; pneumonia; bronchitis; heart disease; sarcoidosis;
 KW sinusitis; purulent otitis media; erythema nodosum; pharyngitis;
 KW vaccine; neutralising epitope.

XX Chlamydia pneumoniae.

XX PN WO9927105-A2.

XX PD 03-JUN-1999.

XX PF 20-NOV-1998; 98WO-IB01890.

XX PR 04-NOV-1998; 98US-0107078.

XX PR 21-NOV-1997; 97FR-0014673.

XX PA (GIST) GENSET.

XX PI Griffais R;

XX DR WPI; 1999-357842/30.

XX PT Genome sequence of Chlamydia pneumoniae

XX PS Page 1170-1171; Disclosure; 1912pp; English.

CC AAV34584-Y35879 represent the proteins encoded by all the open reading
 CC frames in the complete genome (see AAV34584) of Chlamydia pneumoniae.
 CC C. pneumoniae causes respiratory disease such as pneumonia and
 CC bronchitis and is thought to be a contributing factor in heart
 CC disease, sarcoidosis, sinusitis, purulent otitis media, erythema
 CC nodosum or pharyngitis. The polypeptides encoded by the open reading
 CC frames of the C. pneumoniae genome (see AAV34584-Y35879) can be used in
 CC immunogenic compositions as vaccines. Vectors containing C. pneumoniae
 CC nucleotide sequences can also be used as immunogenic compositions,
 CC especially where the vector directs the expression of a neutralising
 CC epitope of C. pneumoniae.

XX SQ Sequence 469 AA;

alignment_scores:

Quality: 450.50 Length: 470
 Ratio: 1.532 Gaps: 16
 Percent Similarity: 62.553 Percent Identity: 29.149

alignment_block:

US-09-303-518D-131 x AAV35375

Align seg 1/1 to: AAV35375 from: 1 to: 469

4 ATTAATAACAAAGAGCTAAATCTGCCCATCGCGGACACCG...GA 50

5 lIleThrValAsnArgGlyLeuAspLeuSerLeuGlnGlySerProLysG 21

51 GCAAGTCATTTATGACGGCCCGCCATTACCGAAGTCGCTTGGCG 100

121 usGlyPheTyrAsn.....LysileAspp 30
 101 AAGAATATGTCGCC.....ATGCGCGCCC
 30 roGluPheValSerIleAspLeuArgProPheGlnProLeuSerLeuLys 46
 127 ATGAAATCAAGAGGTAAGCGCTCAAAAGCCCAAGTCTGTTTGA 176
 47 LeuLysValGluGlnGlyAspAlaValCysSerGlyAlaProIleAlaG 63
 177 AGACAAAAGAAATCGCGCTAGTATTACTCGCGCGCTTCAGCAAAA 226
 63 uTyrLysHisPheProAsnThrTyrIleThrSerHisValSerGlyVal 80
 227 TCGCGCTATTACCGTGGCGAAAGCGCTACTTTCAGTCACTGCTGATT 276
 80 alThrAlaIleArgArgGlyAsnLysArgSerLeuLeuAspValIle 96
 277 ...GCCGTTGAGGACGACGAAATCGAGTTCGAACGCTACGTACCTGA 323
 97 LysLysThrProGlyProThrSerThrGluTyr...ThrTyrAspLeuG 112
 324 AGCGCTGCAAAATTTAGCAGCAGAAAGTCCGCCCAACCTGATTCAAT 373
 112 nThrLeuSerArgSerAspLeuSerGluIlePheLys.....GluA 126
 374 CAGGCTTATGACTCGCTTCGCACCGCTCGTTTCAGCAAAATCCCTGCC 423
 126 snGlyLeuPheAlaLeuIleLysGlnArgProPheAsp...IleProAla 141
 424 GTAGATCCGAG...CGTTTCGCCATCTTCGTCAATCGATGGACACCA 470
 142 lIleProThrGlnThrProArgAspValPheIleAsnLeuAlaAspAsn 158
 471 TCCGCTGGCTGCCGACCTACGGTCATCATC.....A 502
 158 gProPheThrProSerProGluLysHisLeuAlaLeuPheSerSerArg 175
 503 AAGAAGCGCGCAAGACTTCAACCGCGCTGTGTTGTTATGAGCGCGCTG 552
 175 luGluGlyPheTyrValPheValValGlyValArgAlaIleAlaLysLeu 191
 553 ACCGAACGTAAATCCATGTGTGTAAAGCAGCAGCGCCGACACGTGCGTC 602
 192 PheGlyLeuArgProHisIleValPheArgAspArgLeuThrLeuProth 208
 603 TGAATATGCTGCCAATATC...GAACACATGAATTTGGCGCGCCGCATC 649
 208 rGlnGluLeuLysThrIleAlaHisLeuHisThrValSerGlyProPhe 225
 650 CTGCGCGCTTACGTCGACGACCATTCATTCATCGAGCCAGTCGCGCGG 699
 225 roSerGlySerProSerIleHisIleHisSerValAlaProIleThrAsn 241
 700 AATAAA...ACCGTGTGGACCATCAATTATCAAGACGTGATTGCTATCG 746
 242 GluLysGluValValPheThrLeuSerPheGlnAspValLeuThrIleG 258
 747 ACCTTTCTGTAACAGCGCTCTGAATACCGCGCGCTGTTGCTGCTGG 796
 258 yHisLeuPheLeuLysGlyArgIleLeuHisGluGlnValThrAlaLeu 275
 797 CGGCGC.....CTGCAAGTCAACAAACCGCGCTCTTGGCTACCGTTTG 840
 275 laGlyThrAlaLeuLysSerSerLeuArgTyrValIleThrLys 291
 841 GTTGCGAAGGTCTCAACTTACCGCGCGCAATTTGTTGACGCGACAA 890
 292 GlyAlaSerPheSerLeuLeuLeuAsnLeuAsnAspIleSerAspAsn 308
 891 CCGGCTGATTTCGGTTCGGTTCGATTGACCGTTCGATTCACAGGCGCGC 940

```
XX Disclosure; Page 1216; 1755pp; English.
XX
XX AAY37654-y37949 are encoded by open reading frames (ORFs) of the genome
CC of Chlamydia trachomatis (see AAZ01425). The polypeptides can be used as
CC vaccines against Chlamydia trachomatis. Antisense and ribozyme sequences
CC can also be used to control growth of the microorganism. Chlamydia
CC trachomatis is responsible for a large number of diseases, e.g. eye
CC diseases such as conventional trachoma, nonendemic trachoma,
CC paratrachoma, and inclusion conjunctivitis; genital diseases such as
CC nongonococcal urethritis, epididymitis, cervicitis, salpingitis,
CC perihepatitis, bursitis, chondritis; pneumopathy in breast feeding infants;
CC and venereal lymphogranulomatosis. The polypeptides of the invention
CC may be of use in treating these diseases.
XX
XX Sequence 461 AA;
SQ
alignment_scores:
Quality: 438.00 Length: 463
Ratio: 1.490 Gaps: 14
Percent Similarity: 63.499 Percent Identity: 28.078
alignment_block:
US-09-303-518D-131 x AAY37553 ..
Align seg 1/1 to: AAY37553 from: 1 to: 461
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||| : : : : : ||| : : : : : ||| : : : : :
3 lIeValSerArgGlyLeuAlaPheLysSerLeuLysGlyAlaProLysGI 19
: : : : : ||| : : : : : ||| : : : : :
54 AGTCATTATGATCAGGC . . . . . CCGGCCATTACCGAAGTCGGTGC 94
: : : : : ||| : : : : : ||| : : : : :
19 uSerGlyPheCysGlyLysValAspProThrTyrValSerValAspLeu. 35
: : : : : ||| : : : : : ||| : : : : :
95 TTGGCGAGAATATGTTCGGCATCGCCCCTCGATCAAATCAAGGAAGGT 144
: : : : : ||| : : : : : ||| : : : : :
36 ....ArgProPheAlaProLeuProLeuGluValLysValThrProGly 50
: : : : : ||| : : : : : ||| : : : : :
145 GAAGCGCTCAAAAAAGGCCAGTGCTGTTGAAGACAAAAAGATCCGGG 194
||| : : : : : ||| : : : : : ||| : : : : :
51 AspGlnValThrAlaGlySerProLeuAlaGluTyrLysLeuPheSerGI 67
: : : : : ||| : : : : : ||| : : : : :
195 CGTAGTATTTACTGCGCGGCTTCAGCAAAATCGCCGCTATTACCAGTG 244
||| : : : : : ||| : : : : : ||| : : : : :
67 yValPheIleThrSerProValAspGlyGluValGluIleArgG 84
: : : : : ||| : : : : : ||| : : : : :
245 GCGAAAGCGCTACTTCAGTCAGTCGTGATTCGCGTTGAA...GGCAAC 291
||| : : : : : ||| : : : : : ||| : : : : :
84 lyAsnLysArgAlaLeuLeuGluIleValIleLysLysLysProGlyIle 100
: : : : : ||| : : : : : ||| : : : : :
292 GAGCAATCGAGTTTCAGCGCTACGTACTGAGCGCTGGCAAAATGAG 341
||||| : : : : : ||| : : : : : ||| : : : : :
101 SerGlnThrLysPheSer . . . . . TyrAspLeuGlnSerLeuTh 113
: : : : : ||| : : : : : ||| : : : : :
342 CAGCGAAAAGTGGCGCGCAACTGATTCAATCAGGCTTATGAGCTGGC 391
||| : : : : : ||| : : : : : ||| : : : : :
113 rGlnLysAspLeuLeuGluValPheLysLysGluGlyLeuPheAlaLeuP 130
: : : : : ||| : : : : : ||| : : : : :
392 TTCGCACCGCTCGCTTCAGCAAAATCCCTGCGGTAGATGCCGAG...CCG 438
||||| : : : : : ||| : : : : : ||| : : : : :
130 helysGlnArgProPheAsp...lIeProAlaLeuProThrGlnSerPro 145
: : : : : ||| : : : : : ||| : : : : :
439 TTCGCCATCTTCGTCATCGATGTCAGCACCAATCCGCTG.....GCTGC 482
||| : : : : : ||| : : : : : ||| : : : : :
146 ArgAspValPheIleAsnLeuAlaAspAsnArgProPheThrProSerVa 162
: : : : : ||| : : : : : ||| : : : : :
483 CGACCCCTACGCTCATCAAAAGAACGCCGCCGAGAAC.....T 520
||||| : : : : : ||| : : : : : ||| : : : : :
162 lGluLysHisLeuSerLeuPheSerSerLysGluAspGlyTyrTrilleP 179
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521 TCAACGCGGCGCTGTGGTATTGAGCGGCTGACCGAAGCTAAATCCAT 570
```

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179 heValValGlyValGlnAlaLeuAlaLeuPheGlyLeuLysProHis 195
571 GTGTGTAAGCAGCAGCGCAGACGTCGCTGTGAAATATCTGCCAATAT 620
196 IleIleSerThrAspArgLeuThrLeuProThrGlnAspLeuValSerI 212
621 C...GAACACATGAATTTGGCGCGCGCATCTGCCGCTTGAGTGGCA 667
212 eAlaHisLeuHisThrIleAspGlyProPheProSerGlySerProSer 229
668 CGCACATTCATTCATCGACCCAGTC...GGCGCGAATAAAACCGGTGG 714
229 hrHisIleHisIleAlaAArgIleArgAsnGluArgAspValValPhe 245
715 ACCATCAATTAACAGACGCTGATGCTATCGGACGCTTCTGCTAACAGG 764
246 ThrIleSerPheGlnGluValLeuSerIleGlyHisLeuPheLeuLysG 262
765 CGCTCTGAATACCGAGCGGTGCTGCTGCTGGCGGC...CTGCAAG 808
262 yPheValLeuGlyGlnGlnIleValAlaLeuAlaGlySerAlaLeuProp 279
809 TCACAAACCGCGCTCTTCGCTACGTTTGGTGGCGAAGGTGTCTCAA 858
279 roSerGlnArgLysTyrLeuIleThrAlaLysGlyAlaSerPheSerAsp 295
859 CTTACCGCGCGCGAATTTGCTGACGCGGACAAC...CGCGTCAATTCGG 905
296 LeuLeuProLysAspIlePheSerSerAspGluIleThrLeuIleSerG 312
906 TTCGGTATGACGGTGGCATTGCACAGCGCGCATGAT...TATTTGG 952
312 yAspProLeuThrGlyArgLeuCysLysLysGluGluAsnProCysLeu 329
953 GACGCTACCAACATCAGATTCCTGATTCGAGAGAGCGCGCAGCAAGAG 1002
329 lyMetArgAspHisThrIleThrLeuProAsnProLysThrArgGlu 345
1003 CTGTTGCGGTGGTTCGCGCGCAGCGGACAAATCTCCATCAGCGCGCAC 1052
346 SerPheSerPheLeuArgLeuGlyTrpAsnLysLeuThrValThrArgTh 362
1053 CACTCTCGGCCATTCCTTAAACAAA...CTCTTCAAGTTCACGA 1096
362 rTyrLeuSerGlyPhePheLysArgLysArgValPheMetAspMetAsp 379
1097 CAGCGCTCAACGCGCGCGCAGCGCATGGTACCGCATCGGCACATTATG 1146
379 hrAsnMetHisGlyGluLysArgProIleIleAspAlaGluIleTyrGlu 395
1147 CGCGTAATCGCGTGGACATCTGCTACCTTGTCTTTTTCGCGATTTAAT 1196
396 ArgValSerAlaIleProValProValAlaLeuIleIleLysAlaLeuG 412
1197 CGTCGGCATACCGACAGCGCGCGCTTGGTGTGCTGGAATGGACG 1246
412 uThrGlnAsnPheGluGluAlaCysArgLeuGlyLeuGluValAlap 429
1247 AAGAGACCTCGCTTGTGACGCTGCTGCTGCGCGGCAATACGAATAC 1296
429 roGluAspPheAlaLeuProThrPheIleAspProSerLysThrGluMet 445
1297 GGCCCGCTGTTGCCAAGTCTCGAACCATTGAGAG 1335
446 PheSerIleValLysGluSerLeuLeuArgThrGlnLys 458

```

seq_name: /SIDSI/gcgdata/geneseq/geneseq-emb1/AA1999.DAT: AAY34467

seq_documentation_block:

ID AAY34467 standard; Protein; 443 AA.

XX

AC AAY34467;

```

XX 25-AUG-1999 (first entry)
XX Porphyromonas gingivalis protein PG122.
DE Porphyromonas gingivalis; PG; periodontal disease; gingivitis;
XX vaccine; antigenic.
KW Porphyromonas gingivalis.
XX OS
XX Porphyromonas gingivalis.
XX PN WO9929870-A1.
XX 17-JUN-1999.
XX 10-DEC-1998; 98WO-AU01023.
XX 04-AUG-1998; 98AU-0005028.
XX 10-DEC-1997; 97AU-0000839.
XX 31-DEC-1997; 97AU-0001182.
XX 30-JAN-1998; 98AU-0001546.
XX 10-MAR-1998; 98AU-0002264.
XX 09-APR-1998; 98AU-0002911.
XX 23-APR-1998; 98AU-0003128.
XX 05-MAY-1998; 98AU-0003338.
XX 22-MAY-1998; 98AU-0003654.
XX 29-JUL-1998; 98AU-0004917.
XX (CSLC-) CSL LTD.
XX Agius CW, Barr IG, Hocking DM, Margetts MB, Patterson MA;
XX Ross BC, Rothel LJ, Webb EA;
XX WPI; 1999-385613/32.
XX N-PSDB; AAX91685.
XX Antigenic Porphyromonas gingivalis peptides for preventing
XX gingivitis
XX Claim 1; Page 445-446; 588pp; English.
XX AAX91536 to AAX91801 encode two hundred and sixty six antigenic
XX Porphyromonas gingivalis (PG) polypeptide sequences given in AAY34318 to
XX AAX34583. AAX91802 to AAX91989 represent PCR primers used in the
XX isolation of the PG polypeptides. The PG polypeptides have antibacterial
XX activity with a vaccine mechanism of action. The PG polypeptides can be
XX used as vaccines especially against Porphyromonas gingivalis. Probes can
XX be used to detect Porphyromonas gingivalis in standard hybridisation
XX assays. Porphyromonas gingivalis is involved in periodontal disease
XX especially gingivitis.
XX Sequence 443 AA;
XX
XX alignment_scores:
XX Quality: 161.50 Length: 490
XX Ratio: 0.699 Gaps: 22
XX Percent Similarity: 47.143 Percent Identity: 20.612
XX
XX alignment_block:
XX US-09-303-518d-131 x AAY34467 ..
XX
XX Align seg 1/1 to: AAY34467 from: 1 to: 443
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XX 37 GCGGCGCAGACCGGAGCAAGTCATTTATGACGCGCGCCATACCGAAGT 86
XX |||||:||||:||||:||||:||||:||||:||||:||||:||||:
XX 19 AlaGlyLysProValGluValLeu.....ProIleProSerGlnVa 32
XX |||||:||||:||||:||||:||||:||||:||||:||||:||||:
XX 87 CGCGTTGCTGGCGAAGATAATGTCGGCATGCGCCCTCGATGAAATCA 136
XX |||||:||||:||||:||||:||||:||||:||||:||||:||||:
XX 32 lValIleProLeuGlyGlnHisIleGlyAlaProAlaThrAlaThrVal 49
XX |||||:||||:||||:||||:||||:||||:||||:||||:||||:
XX 137 AGGAGGTGAAGCGCTCAAAAAAGCCCAAGTCTGTTTGAAGACAAAAG 186
XX |||||:||||:||||:||||:||||:||||:||||:||||:||||:

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376 gProAlaSerThrValAlaAlaArgArgLysLys.....ProValArg. 390
912 ATTGAACGGTGGCATTGCACAGCGCGCATGATTATTGG..... 952
391LysValAlaAlaGlnSerArgProSerCysTrpLysSerArg 405
953GACGCTACCAACATCAGATTCCGTTATCGAAGAGCCGCGAC 996
406 SerMetThrAlaThrThrGlyArgThrProThrCysAsnSerAlaArgAr 422
997 AAAGAGCTGTTCGGCTGGTGGCGCGGCGGCGGACAAATATCTCCATCAC 1046
422 gProValIleSer.....ArgArgSer.....ProSerA 432
1047 GC.....GCACCACTCTCGGCATTTCTCTAAACAAACACTTCAAGT 1090
432 rgMetPheGlyArgLeuSerAlaSerSerIleAsnMetArgSerThrSer 448
1091 TCACGACACCGCTCAACGCGCGGCGGCGGCGCATGGTACCGCATCGGCAC 1140
449 ValSerAlaProArgThrCys.....ArgAlaThrSe 459
1141 TATGACGCGGTAATGCGGTGGACATCTGCG..... 1171
459 rSerSerAlaSerCysArgCysLeuSerCysProGlnSerThrThrAlaA 476
1172CTACCTGCTTTTGGCGGATTTAATCGTCGCGC 1204
476 laTrpAsnSerGlyTrpThrProAlaProCysProSerSerProMetAla 492
1205 ATACCGACAGCGCGCAGG.....CTTTGGGTGC 1233
493 GlyThrThrArgSerArgArgSerSerArgArgThrProSerTrpProSe 509
1234 TTGGAATTGGAGAAAGACCTCGCTTTGTGCAGCTTCGTCGTCGCGCGG 1283
509 rArgAsnTrpYrSerArgArgArgAsnThrProSerSerAsnSerAlaL 526
1284 CAATACGAATACGCGCGCTGTTCGCAAAAGTCG 1318
526 ysArgArgArgThrGlyLysValSerArgLysCys 537

seq_name: /SIDSL/gcgdata/geneseq/geneseq-emb1/AA2001.DAT: AAB59826

seq_documentation_block:

ID AAB59826 standard; Protein: 1615 AA.

XX AAB59826;

XX 04-APR-2001 (first entry)

XX Protein #3 encoded by TutD/E gene.

XX Toluene degradation; enzyme; waste degradation; TutE; TutD.

XX Thauera aromatica.

OS Xanthomonas maltophilia.

OS Geobacter metallireducens.

OS Azorarcus toluylticus.

XX W0200072650-A2.

XX 07-DEC-2000.

XX 24-MAY-2000; 2000WO-US14298.

XX 01-JUN-1999; 99US-0323872.

XX (UYOH-) UNIV OHIO.

XX Coschigano PW;

XX WPI; 2001-041080/05.

DR N-PSDB; AAF23627.

XX Composition comprising toluene degrading enzyme useful for biological
PT treatment of organic compounds, especially for degrading toluene or its
PT analogs

XX Disclosure; Fig 12; 122pp; English.

XX The present invention relates to toluene degrading enzyme genes and
CC proteins tutH (see AAF23629 and AAB59831), tutI (AAF23630 and AAB59832),
CC tutF (AAF23631 and AAB59833) and tutG (AAF23632 and AAB59834). The
CC toluene degrading enzymes are homologues of pyruvate formate lyase. The
CC toluene degrading enzymes are useful for biological treatment of organic
CC compounds and in particular for the degradation of toluene and its
CC analogs contained in liquid or solid waste source. The present sequence
XX is a protein sequence encoded by toluene degrading enzyme gene, TutD/E.

SQ Sequence 1615 AA;

alignment_scores:

Quality: 145.50 Length: 497

Ratio: 0.677 Gaps: 27

Percent Similarity: 43.260 Percent Identity: 24.748

alignment_block:

US-09-303-518D-131 x AAB59826 ..

Align seg 1/1 to: AAB59826 from: 1 to: 1615

38 CGGCACAGCGGCAAGTCATTTATGACGCGCGCGCCATTACCGAAGTC 87

699 ArgAlaSerSerProLysSerIleSerProLysProArgProThrCy 715

88 GCGT.....TGCTTGGCGAAGATATGCGCATGCGCCCTCGAT 128

715 sArgProSerProGlyThrAlaArgValSerThrThrSerProArg. 731

129 GAAATCAAGGAGGTGACCGCTCAAAAGCCCAAGTCTGTTGAAG 178

732 SerThrThrGlyArg.....ArgTrpSerSerProAla..... 742

179 ACAAAGAATCGGCGTAGTATTACTGCGCGCTTCAGCAAAATC 228

743 .ArgArgSerAlaGlyArgAlaGlyCysAlaArgSerSerA 759

229 CCGCTATTCA...CGTGGCGAAAGCGGTACTTTCAGTCAGTCGTGAT 275

759 rGlyThrSerArgProIleArgSerAlaArgProSerCysSerLysSer 775

276 TGCCGTGGAAGCAACGAGAAATCGAGTTCGAAACGCTACCTACGGAAG 325

776 .ProThrSerValSer.....AlaPheProProS 785

326 CGCTGCAAAATTTAGCAGCGCAAAAGTCGCGCAACCTGATTCAATCA 375

785 erProAlaArgAlaSerArgThrArgCysArgArgAsnSerLeuProSer 801

376 GGCTTATGGACTCGCTTCGCCCGT.....CCGTTTCAGCAAAAT 416

802 SerValThrArgSerSerAlaThrArgAlaAlaThrProArgArgLysTh 818

417 CCCTGCGGTAGATGCGGAGCGGTTCGCCATCTTCGTCA..... 454

818 rProCysCysGlyArg. ThrThrArgProProSerSerThrArgAsnSer 834

455ATCGATGGACACCAATCCGCTGG.....CT 480

835 SerArgAlaThrTrpMetArgTrpAsnSerSerArgTrpAsnValArgPh 851

481 GCGGACCCCTACGTCATCATCAAGAGCCGCCGGAAGACTTCACACGGG 530

851 eProSerMetAlaProAlaSerArgAlaProThrAlaLysSerSerArgg 868

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531 CCTGTTGGTATTGAGCGCCTGACCGCAACGTAATAATCCATGTTGTTAAAG 580
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868 lyArgThrIleCysSerSerPro.....SerAlaAlaProThr 881
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581 CAGCAGCGCAGAGCTGCGCTCTGAAATGCTGCCAATATCGAAACACAT 630
  ::::: ||| |||
882 ProArgAlaArgThrProAlaThrThrProThrProSerSerArgGlnPr 898
  ::::: ||| |||
631 GAATTTGGCGCGCGC.....ATCCTGCGCGCTT 659
  :: ::::: ||| |||
898 oSerGlySerAlaArgProSerProProSerSerSerAlaIleProArg 915
  :: ::::: ||| |||
660 GAGTGGCAGCCACA.....TTCATTTCATCGAGC..... 688
  ::::: ||| |||
915 rgThrAlaAlaArgArgCysAlaGlyPheSerSerAlaAlaThrAsp 931
  ::::: ||| |||
689 .....CAGTCGGCGCGCAATAAAACCGTG 711
  ::::: ||| |||
932 SerAlaIleArgArgSerSerThrThrArgSerAlaArgSerArgAs 948
  ::::: ||| |||
712 TGGACCATCAATTATCAAGAGCTGATTGCTATCGAGCTTGTTCGTAAC 761
  ||||| ::::: |||
948 nThrProSerSerAlaSerThr..... 955
  ||||| ::::: |||
762 AGCGCGTCTGAATACCGAGCGCGTGGTTCCTTGGCGCGCTGCAAGTCA 811
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956 .....AlaThrAlaPro.....ProThrArgLysPro 964
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812 ACAAACCGCGCTCTGCGTACCCTTTTGGGTGCGAAGGTCTCTCAACTT 861
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965 ThrThrGlySerThrCys.....CysAlaCysAr 974
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862 ACCGCGCGCAATTGGTGTGACCGGACACCGCGTGATTTCCGGTTCGGT 911
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974 gProAlaSerThrValAlaAlaAlaArgArgLys.....ProValArg. 988
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912 ATTGAACGCTGCGATTGCACAAAGCGCGCATGATTATTGG..... 952
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989 .....LysValAlaAlaAlaAlaSerSerArgProSerCysTrpLysSerArg 1003
  ||| ::::: ||| |||
993 .....GACGTACCAATCAGATTTCCTTATCGAAGAGCGCGCAGC 996
  ||||| ||| ||| ::::: |||
1004 SerMetThrAlaThrThrGlyArgThrProThrCysAsnSerAlaArgAr 1020
  ||| ||| ::::: ||| |||
997 AAAGAGCTGTTCGGTGGTTCGCCCGCAGCGGACAAATCTCCATCAC 1046
  ||| ||| ::::: ||| |||
1020 gProValIleSer.....ArgArgSer.....ProSerA, 1030
  ||| ||| ::::: ||| |||
1047 GC.....GCACCACTCTCGGCAATTTCCTAAACAACTCTCTCAAGT 1090
  || ::::: ||| |||
1030 rgMetPheGlyArgLeuSerAlaSerSerIleAsnMetArgSerThrSer 1046
  || ::::: ||| |||
1091 TCACGACAGCCGCTCAACGCGCGCGACCGCGCATGGTACCGATCGGCAC 1140
  ||| ||| ::::: |||
1047 ValSerAlaProArgThrCys.....ArgAlaThrSe 1057
  ||| ||| ::::: |||
1141 TATGACCGGTAATCGCTGGACATCTGC..... 1171
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1057 rSerSerAlaSerCysA-gCysLeuSerCysProGlnSerThrThrAlaA 1074
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1172 .....CTACCTTCCTTTTGGCGGATTAAATCGTCGCGC 1204
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1074 laTrpAsnSerGlyTrpThrProAlaProCysProSerProMetAla 1090
  ||||| ||| |||
1205 ATACGACAGCGCGGAGG.....CTTTGGGTTC 1233
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1091 GlyThrThrArgSerArgSerSerArgArgThrProSerTrpProSe 1107
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1234 TTGGAATTCGAGGAAGAACCTCGCTTTGTGCAGCTTCGCTCGCCCGG 1283
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1107 rArgAsnTrpTySerArgArgArgAsnThrProSerSerAsnSerAlaL 1124
  : ||||| ::::: ||| |||
```

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132 AATCAAGGAAGTGAAGCCG..... 151
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650 spAlaArgAsnArgAspProLeuGlyArgGlyCysThrAspProCysCys 666
152 .....TCAAAAAGCCCAAGTCGCTTTTGAAGACAAAGA... 187
   :::::
667 ProProAlaThrArgArgGlyArgAsnCysCysArgAlaAlaArgAr 683
188 .....ATCCGGGCGTAGTATTACTCGCGCGCTTCAGGCAA 224
   :::::
683 gPheArgGlyProLeuArgAlaAlaProAlaCysAlaArgPheAla 699
225 AATCCCGCTATTCCACG..... 242
   :::::
700 TyrArgArgLeuGlyProArgCysThrSerArgGlyArgSerArgCysSe 716
243 .....TGCGAAAGCCGCTACTTCAGTCAGTCGTG 273
716 rProAspArgCysCysArgTrpSerArgCysSerProSerProArgA 733
274 ATTGC..... 278
733 rGysProProSerSerProAlaGlyAlaProGlyAlaThrCysSerArg 749
279 .....CGTTGAAGCAA.....CGACGAATCGAGT 304
   :::::
750 ArgProPheSerArgSerArgAspSerAlaGlyProArgAlaAlaArgph 766
305 TCGAAGCGTAGCTACCTCAACCGCTGGCAAAATTGACGAGCGAAAAGTG 354
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766 eArgArgCysArg.....AspAlaCysGluArgArgAlaArgC 779
355 CGCGCAACCTGATTCAATCAGGCTTATGGACTGCGCTTCGCACCCGCTCC 404
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779 ysProGlyPro.....ArgSerAlaProSer 787
405 GTTCAGCAAAATCCTCGCTAGATCGCGAGCGCTTCGC..... 443
788 IleArgArgGlySerArgAspArgSerArgAlaSerArgSerArgGlyse 804
444 .....CATCTTCGTCANTCGC 459
804 rProLeuCysGlyAlaThrAlaThrSerCysProArgArgArgCysS 821
460 ATGACACCAATCGCTGGCTGC.....CGACCC 488
   :::::
821 erIleGlyAlaSerSerGlyCysProHisProProValArgArgSerPro 837
489 TACGTCATCATCAAGACCGCCGAGACTTCAACGCGGCTGTGG 538
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838 ValAsnSerSerLysArgAlaHisArgArgCysThrAlaArgArgGlyAr 854
539 TATTGACCCGCT.....GACCGAAGCTAAATCCATGTGTGTA 579
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854 gPheArgGlyProThrSerArgAspThrGlyArgArgCysTrpArgT 871
580 GCACGACGCGCAGCTGCGCTGAAATGCTGCCAATATCGAAACACA 629
   :::::
871 rpProArgProArgArgCys.....ArgCysSerArgArgTrpGlyArg 885
630 TGAATTGGCGCCGCTGCTCGCGCTGAGTGGCCACCATTCATT 679
   :::::
886 ProLeuTrpAlaSerGlyCysProArgAlaArgTrp..... 897
680 TCATCGACCGCAGTCGGCGGAATAAACCGCTGTGGACCATCAATTATCAA 729
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898 .ArgArgGlySerAsnTrpSerSerGlyArgSerSerAlaAlaSerProL 914
730 GAGCTGATTCTATCGGACCTTTGTCGTAAACAGCGCTCTGAATACCGA 779
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914 ysArgThrCysGlyArgArg...ValArgSerAspThrSerAlaArgArg 929
780 GCGCGT..... 785
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930 SerArgCysProAlaSerSerProIleArgTrpThrGlyArgCysArgAr 946
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786 .....GGTTGCCTTGGCGGCTGCAAGTCAACAAACCG 819
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946 gTrpArgArgProLeuGlyCysSerProArg.....A 957
820 CGCCTCTTGCTACCGTTTGGTGGCAAGGTGCTCAACTTAC..... 863
   :::::
957 laThrCysThrAlaArgCysGlyArgAspGlyCysSerAlaPheGly 973
863 ..... 863
974 AsnProLeuHisArgSerLeuArgGlyProTrpAlaAlaProPheArgAl 990
864 .....CGCGCGCAATTGGTTGACGCGACACACCGC 894
   :::::
990 aHisArgSerArgSerThrThrArgArgCysAlaValArgGlySerSerA 1007
895 GTGATTTCGGTTCGGTATTGAACGGTGCATTGCACAAG..... 934
   :::::
1007 rHisAspArgThrAlaSerThrArg..ArgProHisLysProProLysG 1023
935 .....GCGCGCATGATTATTGGGACGCTACCACA 964
   :::::
1023 lyCysAlaThrAspIleHisSerGlyArgTyrcysTrpProArgThrAla 1039
965 ATCAGATTTCGGTATCGAAGAGCGCGCAGCAAGAGCTGTTCGGCTGG 1014
   :::::
1040 .....SerSerArgAlaAlaSerGlyAlaSerAlaLysAr 1051
1015 GTTGGCGCGCAGCGACAAATACTCCATCAGCGCCACCA..... 1054
   :::::
1051 gThrArgLeuArgArgArgSerCysProValArgSerProArgArgArgG 1068
1055 ..CTCTCGGCCATTCTCTAAAAACAACACTTCAAGTTTCACGACGCG 1102
   :::::
1068 lyThrArgAlaAlaThrHisSerAlaCysGlySerSerSerArgArgPro 1084
1103 TCACGCGCGCGCAGCGCCGCTATGG.....TACCGATCGGCATTA 1143
   :::::
1085 SerSerGly.....ArgProTrpSerValProIleArgProSerSerIl 1099
1144 GAGCGCGTAAATCGCTTGA..... 1163
   :::::
1099 eCysGly.ArgAlaValGlyLeuThrSerProSerProLeuAsnArg 1115
1164 .....CATCCTGCCTACCTTGTCTTTTGGCGCGATTAA 1195
   :::::
1116 ProPheAlaArgArgSerAlaProAlaSerThrProCysArgArgHisA 1132
1196 TCGTCGCGGATACCGACGCGCGAGCTTGGTGTGCTTGAATTGAC 1245
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1132 nArgArgArgTrpGlySerArgArgProPhe.....A 1143
1246 GAAGAAGA 1253
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1143 rgArgArg 1145
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seq_name: /SIDSL/gcgdata/geneseq/geneseq-emb1/AA2001.DAT:AA59817

seq_documentation_block:

ID AAB59817 standard; Protein; 999 AA.

XX AAB59817;

XX

XX

DT 04-APR-2001 (first entry)

XX

DE TutD protein #8.

XX

KW Toluene degradation; enzyme; waste degradation; TutD.

XX

OS Thauera aromatica.

OS Xanthomonas maltophilia.
 OS Geobacter metallireducens.
 XX Azarcus toluilyticus.
 PN WO200072650-A2.
 XX 07-DEC-2000.
 XX
 XX 24-MAY-2000; 2000WO-US14298.
 PF
 XX 01-JUN-1999; 99US-0323872.
 XX
 XX (UYOH-) UNIV OHIO.
 PA
 XX Coschigano PW;
 PI
 XX WPI; 2001-041080/05.
 DR N-PSDB; AAF23625, AAF23627.
 DR
 XX
 XX Composition comprising toluene degrading enzyme useful for biological
 PT treatment of organic compounds, especially for degrading toluene or its
 PT analogs -
 XX
 XX Disclosure; Fig 5; 122pp; English.
 PS
 XX The present invention relates to toluene degrading enzyme genes and
 CC proteins tutH (see AAF23629 and AAB59831), tutI (AAF23630 and AAB59832),
 CC tutF (AAF23631 and AAB59833) and tutG (AAF23632 and AAB59834). The
 CC toluene degrading enzymes are homologues of pyruvate formate lyase. The
 CC toluene degrading enzymes are useful for biological treatment of organic
 CC compounds and in particular for the degradation of toluene and its
 CC analogs contained in liquid or solid waste source. The present sequence
 CC is a protein sequence for toluene degrading enzyme, TutD.
 XX
 XX Sequence 999 AA;

alignment_scores:
 Quality: 129.00 Length: 573
 Ratio: 0.617 Gaps: 27
 Percent Similarity: 36.475 Percent Identity: 22.339

alignment_block:

US-09-303-518d-131 x AAB59817 ..

Align seg 1/1 to: AAB59817 from: 1 to: 999

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32 CCATCGCGGGGAGCGGAGCAAGTCAATTAATGACGGCGCGCCATTACC 81
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25 ProSerGlyAlaSerArgSerSerAlaAlaAlaProArgArgProPr 41
82 GAAGTCGGTTCCTGGCGAAGATATGCGCATGCGCCCTCGATGAA 131
|::: ||||| ||||| ||||| ||||| ||||| ||||| |||||
41 oSerHisSerSerAlaAlaTyrGlyAlaSerCysThr.....A,55

132 AATCAAGGAGGTGAAGCG..... 151
:::||||| |||||
55 spAlaArgAsnArgAspProLeuGlyArgGlyCysThrAspProCys 71
152 .....TCAAAAAGCCCAAGTGTGTTGAAGCAAAAAGA.. 187
72 ProProAlaThrArgArgLysArgAsnCysCysArgAlaAlaArg 88
188 .....ATCCGGCGGTAGTATTACTGCGCGCGCTTCAGGCAA 224
:::||||| ||||| ||||| ||||| ||||| ||||| |||||
88 gPheArgGlyProLeuArgAlaAlaProAlaCysAlaAlaArgPhe 104
225 AATCGCCGCTATTACCG..... 242
||||| |||||
105 TyrArgArgLeuGlyProArgCysThrSerArgGlyArgSerArgCys 121
243 .....TGCGGAAAGCGCGTACTTCACTCAGTCAGTCGTG 273
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121 rProAspArgCysCysArgTrpSerArgCysSerSerProSerProArg 138
274 ATTGC..... 278
138 rgCysProProSerSerProAlaGlyAlaProGlyAlaThrCysSerArg 154
279 .....CGTTGAAGCAA.....CGACGAAATCGAGT 304
155 ArgProPheSerArgSerArgAspSerAlaGlyProArgAlaAlaArgph 171
305 TCGAAGCGCTACGTACCTGAAGCGCTGCGCAAAATTCAGCAGCGGAAAGTG 354
171 eArgArgCysArg.....AspAlaCysGluArgArgAlaArgC 184
355 CGCGCGCAACCTGATCAATCAGGCTTATGGACTTGGCTTCGCACCGCTCC 404
184 ysProGlyPro.....ArgSerAlaProSer 192
405 GTTCAGCAAAATCCCTGCGTAGATGCCGAGCGCTTCGC..... 443
193 IleArgArgGlySerArgAspArgSerArgAlaSerArgSerArgSerAr 209
444 .....CATCTTCGTC 453
209 gglySerProLeuCysGlyAlaThrAlaThrSerCysProArgArgArg 226
454 AATGCGATGGACACCAATCCGCTGGCTGC..... 482
226 rgCysSerIleGlyAlaSerSerGlyCysProHisProProValArgArg 242
483 CGACCCCTAGCGTCATCATCAAGAGCGCGCGGAGACTTCAACCGCGCC 532
243 SerProValAsnSerSerLysArgAlaHisArgArgCysThrAlaArgAr 259
533 TGTGTGTTATGAGCGCCT.....GACGGAACGTAAATCCATCTGTG 573
259 gglyArgPheArgGlyProThrSerArgAspThrGlyArgArgArgCysT 276
574 TGTAAAGCAGCAGCGCGACGCTGCTGAAATGCTGCCAATATCGA 623
276 rpArgTrpProArgProArgArgCys.....ArgCysSerArgArgTrp 290
624 AACACATGAATTTGGGCGCGCATCTCTCGCGCTTGAGTGGCAGCAGCA 673
291 GlyArgProLeuTrpAlaSerGlyCysProArgAlaArgTrp..... 304
674 TTCATTTTCATCGACGCGCGGCAATAAAACCGTGTGGACCATCAAT 723
305 .....ArgArgGlySerAsnTrpSerSerGlyArgSerSerAlaAla 319
724 TATCAAGACGTGATTGCTATCGACGCTTTGTCGTAACAGCGCGCTCAA 773
319 erProLysArgThrCysGlyArgArg...ValArgSerAspThrSerAla 334
774 TACCGACGCGGT..... 785
335 ArgArgSerArgCysProAlaSerSerProIleArgTrpThrGlyArgCy 351
786 .....GGTTGCTTGGGCGCGCTGCAAGTCAAC 813
351 sArgArgTrpArgProLeuGlyCysSerProArg..... 363
814 AAACCGCGCTCTGCGTACCGTTCGTCGAGAGGTGCTCAACTTAC 863
364 .....AlaThrCysThrAlaArgCysGlyArgAspGlyCysSerAlaPhe 378
863 ..... 863
379 PheGlyAsnProLeuHisArgSerLeuArgGlyProTrpAlaAlaProPh 395
864 .....CGCGGCGAATTTGTTGACGCGGAC 888
395 eArgAlaHisArgSerArgSerThrThrArgCysAlaValArgGlyS 412

```


XX Mycobacterial DNA vectors containing reporter constructs - for
PT identifying coding or promoter sequences involved in
PT infection-associated protein expression
XX
XX Claim 32; Fig 41b; 309pp; French.
XX
XX Sequences AAY04742-Y05000 and AAY07201-Y07204 represent secreted
CC proteins from various Mycobacterium species microorganisms. The
CC encoding nucleotide sequences can be used as primers and probes for
CC methods for detecting and identifying mycobacteria, especially belonging
CC to the M. tuberculosis complex. The encoded proteins can be used in
CC vaccines for immunisation against a bacterial or viral infection.
XX
SQ Sequence 573 AA;

alignment_scores:
Quality: 119.00 Length: 477
Ratio: 0.626 Gaps: 20
Percent Similarity: 39.832 Percent Identity: 23.270

alignment_block:

US-09-303-518D-131 x AAY04955 ..

Align seg 1/1 to: AAY04955 from: 1 to: 573

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166 GlyAsnTyrArgLeuGlyAlaAlaGlyArgArgSerArgArgProVa 182
122 CCTCGATCAAAATCAAGAGGTGAACCGTCAAAAAGG..... 161
182 lArgAlaArgGlyValGlyArgCysGlyHisArgArgArg***ArgGlyG 199
162CCAAAGTCTGTTTGAAGACAAAGAAATCC 191
199 lHisArgAlaGlyLysAspProArgThrAla***ArgAlaArgArgCys 215
192 GGGCGTAGTATTACTGGCGCGGCTTCAGCAAAATCGCGCTATTCCAC 241
216 GlyArgGly.....GlyArgArgArgThrGlyPr 225
242 GTGGCGAAAGCGGTACTTCACTCAGTCGTCGATTCGCGTTGAAGCAAC 291
225 oAlaGlySerAlaGlyArgValAlaLeuHisHisLeuArgAlaGlyThrC 242
292 GACGAATCGAGTTCGAACGCTACGTACCTGAAGCGTGGCAAAATTCAG 341
242 ysProGlyGlyLeuArgThrLeu.....AlaAlaArgValAla 254
342 CAGCGAAAAGTGGCGCGCAACCTGAT.....TC 370
255 AspArgHisGlyTyrProThrProArgProAlaLeuArgGlyAspValSe 271
371 AATCAGGCTTATGGACTCGCTTCGACCCGTCGCTTCAGCAAAATCCCT 420
271 rValGlyGlyMet***CysCysSerGlyGlyProVal...AlaGlySerT 287
421 GCCGTAGATGGCGGCGGCTTCGCTTCGCTTCGCTTCGCTTCGCTTCG 470
287 hrGlnGlyIleGly***ValGlyGlyHisArgArgCysSerAlaArgGln 303
471 TCCGCTGGCTCGCGACCTACGGTATCATCAAGAAAGCGCGCAAGACT 520
304 LeuLeuArgThrArgPro.....HisArgArgArgArgCysArgArgG 318
521 TCAAAACGGCGGCTGTGTATGTAGCCGCTGACGCAACGTAATCCAT 570
318 ySerArgIleGlyGlyAlaSer***ProAspArgAspLeuGly.... 333
571 GTGTGTAAAGCAGCAGCGCAGACGTGCGCTGAAATGCTGCCAATAT 620

334AlaArgPhe 336
621 CGAAACACATGAATT.....TGCGGGCCCATCTCTGCGGCTTGA 661
337 ArgAspGlnArgIleAlaGlyArgTrpLeuAspAlaGlyProArg..... 351
662 GTGGCAGCACATTCATTTCATCGAGCGAGTCGGCGGAATAAAACCGTG 711
352ArgAlaGlyGlyArg..... 357
712 TGGACCATCAATATCAAGACGTGATTGCTATCGGACGTTTGTTCGTAAC 761
358ArgArgArgCysArgArgAla..... 364
762 AGGCGCTCTGATACCGAGCGGTGGTTCCTTGGCGGCTGCAAGTCA 811
365ValArgArgGlyArgLeuArgAlaAlaThrGlySe 377
812 ACAAAACCGCGCTCTTGGTACCGCTTTTGGGTGC.....GAAG 849
377 rArgArgArgAspThrGlyArgArgTyrGlnCysProProAlaGlyAlaG 394
850 GTGTCTCAACATTACCGCGCGGCAATTGTTGACCGGACACCGCGGTAT 899
394 lArgGlyArgHisArgArgAla..... 402
900 TTCCGCTTGGGTATTGAACGGTGCATTCACAGCGCGCATGATTATT 949
403ArgAspGlyAlaAlaGlnTrpLeuCysGlyArgArgArg...ThrGl 417
950 TGGGACGTTACCAATCAATCAGATTTCGTTATCGAAGAGCGCGCAAAA 999
417 yGlyArgValTyrArgGlyAsp...ArgLeuGlyArgArgArgGlyThrA 433
1000 GAGCTTTCGCTGGTTCGCGCGCGGCAAGCAATA..... 1037
433 rGAlaAspArgIleAspGlyAlaGlyValGlyArgAlaGlyArgAla*** 449
1038CTCCATCAGCGCGCACCATCTCG 1060
450 ArgGlyProProGlyArgArgArgLeuGlnHisGlyProCysArgArg 466
1061 GCCATTTCTAAAAACAACCTCTCAAGTTCCAGCAGC..... 1100
466 gCysPheProAlaAlaArgIleGlyAlaHisCysHisProLysGlyAlaAspL 483
1101CGTCAACGGCGCGCGCCAT 1124
483 euGlyArgTyrLeuGlnAlaGlyArgArgSerGlyTyrArgGlyArgArg 499
1125 GSTACGATCGGACCTTATGAGCGGTAATGCGGTGGACATCTGCTGCTA 1174
500 GlyAlaAspArg..... 503
1175 CTTTGTCTTTCGCGGATTTAATGTCGCGCATACCGACAG.....C 1215
504ArgArgArgCysArgArgGlyGlyHisA 513
1216 GCGCAGCGCTTGGGTTGCTTGGAAATGGAGAAAGACCTCGCTTGTG 1265
513 rgSerGlyArgProValValGlyIleGlyArgArgSerGlyAspGlyAla 529
1266 CAGCTTCGT.....CTGCCCGGCAATACG 1291
530 AsnTrpArgArgArgAsnArgArgArgGlyCysArgProGlyThrAlaCy 546
1292 AATACGCGCGCTGTTCCGCAAGTCTGGA 1322
546 sAlaArgProProSerArgHisArgAlaGly 556

seq_name: /SIDS1/gcgdata/geneseq/geneseq-emb1/AA2000.DAT: AAB15935
seq_documentation_block:

ID AAB15935 standard; Protein; 740 AA.

XX AAB15935;

XX 05-OCT-2000 (first entry)

XX E. coli proliferation associated protein sequence SEQ ID NO:292.

XX Escherichia coli; E. coli; proliferation; inhibition; screening;

KW antimicrobial; bacterial growth; antisense therapy; antibacterial.

XX Escherichia coli.

OS WO200044906-A2.

XX 03-AUG-2000.

XX 27-JAN-2000; 2000WO-US02200.

XX 27-JAN-1999; 99US-0117405.

XX (ELIT-) ELITRA PHARM INC.

XX Zyskind J, Ohlsen KL, Trawick J, Forsyth RA, Froelich JM, Carr GJ;

PI Yamamoto RT, Xu HH;

XX WPI; 2000-514822/46.

XX N-PSDB; AAB65940.

XX Novel polynucleotides and polypeptides associated with microorganism

PT proliferation, used to identify inhibitors of bacterial growth and

PT proliferation, for use in antisense therapy.

XX Claim 11; Page 217-219; 316pp; English.

XX AAA65809 to AAA65889 and AAA66058 to AAA66138 represent nucleotide
sequences derived from Escherichia coli which inhibit E. coli
proliferation. AAA65890 to AAA66055 and AAB15886 to AAB16040 represent
nucleotide and protein sequences associated with E. coli proliferation.
XX AAA66056 and AAA66057 represent primers used for sequencing E. coli
proliferation inhibiting nucleotide inserts in an example from the
present invention. Methods from the present invention can be used to
identify a proliferation-required gene in a microorganism, by contacting
a microorganism with a proliferation-required gene activity inhibitory
nucleic acid identified in another organism, and determining if
inhibition occurs in the second microorganism. The nucleic acid sequences
identified as being required for bacterial growth and proliferation, can
be used for antisense therapy for killing bacteria.

XX Sequence 740 AA;

alignment_scores:

Quality	Length
117.00	313
Ratio: 0.731	Gaps: 13
Percent Similarity: 51.118	Percent Identity: 22.684

alignment_block:

US-09-303-518D-131 x AAB15935 ..

Align seg 1/1 to: AAB15935 from: 1 to: 740

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48 LysGlnHisIleGlyAlaGluGlyLeuGlyCysValSerValGlyAspLeu 64

150 CGTCAAAAAGCGCAAGTGTGTTGAAGACAAAAGAAATCCGGCGTAG 199

64 sValLeuArgGlyGlnProLeuThrArgGlyArgGlyLysMetLeuProV 81

200 TATTACTCGCGCGCTTCAGGAAATCCCGCGTATT 237

81 alHis....AlaProThrSerGlyThrValThrAlaIleAlaProHisSer 96

238CACCGTGGCGGAAAAGCGCGTACTTTCAGTCAGTCGTGATGCGGT 281
97 ThrAlaHisProSerAlaLeuAlaGluLeu...SerValIleIleAspAl 112
282 TGAAGGCAACGACGAAATCGAGTTTCGACGCTAGTACCTGAA..... 324
112 aaspGlyluasp.....CystripileProargAspGlyT 124
325GCCTGGCAAAATTCAGACGCGGAAAGTCGCGCGCAACCTGATT 369
124 rpAlaAspTyrArgThrArgSerArgGluGluLeuIleGluArgIleHis 140
370 CAATCAGCGTTATGAGTCGCGCTTCGACACCGCTCCGTTTCAGC..... 411
141 GlnPheGlyVal...AlaGlyLeuGlyAlaGlyPheProThrGlyVa 156
412 .AAATCCCTGCGTAGATGCCGAGCGGTTCCGCATCTTCGTCATCGCAATCGGA 460
156 lLysLeuGlnGlyGlyAlaGlyIleGluThrLeuIleIleAsnAlaA 173
461 TGGACACCAATCCGCTGCGTCCGCGACCCCTACGGTTCATCATCAAGAGGCC 510
173 laGluCysGluProTyrIleThrAlaAspArgLeuMetGlnAspCys 189
511 GCCGAGAGACTTCAACGCGCGCTGTTGGTATTGAGCGCGCTGACCGAA.. 558
190 AlaAlaGlnValValGluGlyIleArgIleLeuAlaHisIleLeuGlnPr 206
559CGTAAATCCATG 571
206 oArgGluIleLeuIleGlyIleGluAspAsnLysProGlnAlaIleSerM 223
572 TGCTGAAGCAGGCGGCGACGTCGCGCTCTGAAATGCTGCCAATATC 621
223 etLeuArgAlaValLeuAlaAsp.....SerAsnAspIle 234
622 GAAACACCAATTTGGCGCGCGCATCTCCGCGGC..... 657
235 SerLeuArgValIleProThrLysTyrProSerGlyGlyAlaLysGlnLe 251
658TTGAGTGGCAGCAGCATTCATTTTCATCGAGCCAGTCGCGG 697
251 uThrTyrIleLeuThrGlyLysGlnVal.....ProHisGlyG 264
698 CGAATAAACCGTGGACCATCAATATCAAGACGTG.....ATT 738
264 lYArgSerSerAspIleGlyValLeuMetGlnAsnValGlyThrAlaTyr 280
739 GCTATCGGACGCTTTGTTGTAACAGCGCGTCTGAATACCGAGCGGTGT 788
281 AlaValLysArgAlaValIleAspGlyGluProIleThrGluArgValVa 297
789 TGCCTTGGCGGCGCTCAAGTCAACAAACGCGCGCTCTTCGTCACCGTTT 838
297 lThrLeuThrGlyGluAlaIleAlaArgProGlyAsnValTrpAlaArgL 314
839 TGGTGTGCGAAGGTGCTCAACTT.....ACGCGCGCGCAATTGTTGAC 882
314 euGlyThrProValArgHisLeuLeuAsnAspAlaGlyPheCysProSer 330
883 GCGGACACCGCGGTATTTCGCGTTCCGGTTTCGGTATTGTAACCGGT 921
331 AlaAspGlnMetValIleMetGlyGlyProLeuMetGly 343

seq_name: /SIDS1/gcgdata/geneseq/geneseq-emb1/AA2001.DAT:ABG03731

seq_documentation_block:

ID ABG03731 standard; Protein; 696 AA.

XX ABG03731;

XX 13-FEB-2002 (first entry)


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847 isAsnLysValAlaAspIleGluLysLysAlaArgGluLeuArgAspLeu 863
499 ATCAAGAAGCCCGGAGAGCTTCAACGCGCGCTTGGTATTGAGCGG 548
864 ValProGluAlaArgValValValAlaHisGlyGlnMetSerGluGlu 880
549 CCTG.....ACCGAACGTAATAATCCATG 571
880 uLeuGluGlnThrValGlnGlyPheTrpAspArgGluTyrAspValLeu 897
572 TGTGTAAGCA.....GCAGCGCCAGACGTGCGCTCGTCAAAATGCT 612
897 alCysThrThrIleValGluThrGlyLeuAspIleSerAsnAlaAsnThr 913
613 GCCAATATCGAAACACATGAATTTGGCGCCCGCATCTCCGCGGCTTGAG 662
914 LeuIleValGlu.....AsnAlaHisHisMetGlyLeuSe 925
563 TGGCAGCAC..... 672
925 rGlnLeuHisGlnLeuArgGlyArgValGlyArgSerArgGluArgGlyT 942
673 ..ATTCAATTCATCGAGCGAGTGGCGCGAATAAAACCGTGTGACCATC 720
942 yAlaTyrPheLeuTyrProLysGlyAla.....ThrLeuThrGluMet 956
721 AATTATCAAGCGTATGCTATCGGACGTTTGTTCGTACACAGCGCTCT 770
957 SerTyrAspArgLeuAlaThrIleAlaGln.....AsnAsnAspLe 970
771 GAATACCGAGCGGTGCTGCTTGGCGCGCTGCAAGTCAACAAACCGC 820
970 uGlyAlaGlyMetAlaValAlaMetLysAspLeuGluMet.....A 984
821 GCCTCTTCGTACCGTTTGGTGGCGAGGTGCTCAACTACCGCC... 867
984 rGlyAlaGlyAsnValLeuGlyAlaGluGlnSerGlyHisIleAlaGly 1000
868 .....GGCAATTGCTTACGCGGA 887
1001 ValGlyPheAspLeuTyrValArgLeuValGlyGluAlaValGluAlaTyr 1017
888 CAACCGGTGATTTCGGTTCGTTGAACGGTGGATT..... 927
1017 rArgAlaLeuAlaAspGlyLysValValAspGlyThrValLysGlyProL 1034
928 .....GCACAAGCGCGCATATTAT 948
1034 ysGluIleArgValAspLeuProValAspAlaHisIleProGluLysTyr 1050
949 TTGGGACGTACCAACATCAGATTTCGTTATCGAAGAGCGCCGACAA 998
1051 Ile..AsnAlaGluArgLeuArgLeuGluIleTyrArgLysLeuAlaGlnS 1067
999 AGAGCTGTTCGGTGGTGGCGCGAGCGCGCAATAATCTCCATCAGCG 1048
1067 erGluSerGluValAspLeuArg..LeuAlaValGluGluMetGluAspAr 1083
1049 GCACCACTCTCGGCATCTTCCTAAACAACTCTCACTTCCACGACA 1098
1083 gTyrGlyProIleProGluGluValGluArgLeuLeuAlaValSerArgL 1100
1099 GCGGTCAACGCGCGCGACCGCGCTGTTAC.....GATCGG 1136
1100 euArgHisLeuMetArgGluAlaHisLeuThrAspIleAlaValGln..G 1116
1137 CACTTATGAGCGGTAACTGGTGGACATCTCGCTACC.....TTGC 1180
1116 yThrArgIleLysValHisProValAspLeuAlaAspSerGlnGlnValA 1133
1181 TTTTCGCGATTATCTCGTCGATACCGACGCGCGAGCTTTGGGT 1230
1133 rgLeuLysArgLeuPheProGlyAlaThrTyrArgAla....AlaAlaLys 1148
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1231 TGCTTGAATTGGACGAAGAAGACCTCGCTTGTGACAGCTTCGTCGCC 1280
1149 AlaIleGlnLeu.....SerPheProLysTh 1157
1281 GGCAATATCAGAAATACGCGCGCTGTTCGCGCAAGTG..... 1317
1157 rGlyAsnLysValThrAspProLeuLeuArgAspValAspLeuGlnT 1174
1318 .....CTGGAACCATTCAGCAAG 1335
1174 rpValAlaAsnPheIleSerAsnMetPheAsnLeuGluGluIleAspVal 1190
1336 GAAGGC 1341
1191 ArgGly 1192
seq_name: /SDSL/gcgdata/geneseq/geneseq-emb1/AA2001.DAT: AAB96063
seq_documentation_block:
ID AAB96063 standard; Protein; 822 AA.
XX AAB96063;
XX AC
XX XX
DT 29-OCT-2001 (first entry)
XX XX
DE Putative P. abyssi pyruvate kinase #1.
XX XX
KW Hyperthermophilic archaeon; hyperthermophilic protein.
XX OS
XX PYROCOCCLUS ABYSSI.
XX FR2792651-A1.
XX PN
XX XX
PD 27-OCT-2000.
XX XX
PF 21-APR-1999; 99FR-0005034.
XX XX
PR 21-APR-1999; 99FR-0005034.
XX XX
PA (CNRS ) CNRS CENT NAT RECH SCI.
PA (IFRE-) IFREMER INST FR RECH EXPL MER.
PI Forterre P, Thierry JC, Prieur D, Dietrich J, Lecompte O;
PI Querellou J, Weissenbach J, Saurin W, Heilig R;
XX WPI; 2001-126236/14.
XX XX
XX New nucleotide sequences isolated from Pyrococcus abyssi encode
XX proteins useful in industry -
XX PS Claim 7; Pages 677-680; 1657pp; French.
XX XX
XX The present invention relates to the genomic sequence of Pyrococcus
XX abyssi (see AAF86431 and AAH41223-7) and P. abyssi proteins. P. abyssi is
XX a hyperthermophilic archaeon, which is isolated from deep-sea
XX hydrothermal vents. The present sequence is one such P. abyssi protein.
XX The proteins of the present invention have various potential industrial
XX uses, since the proteins are stable at very high temperatures, some up to
XX 110 degrees centigrade.
XX Note: This patent is in the same patent family as WO2000065062, which
XX contains additional sequences as shown in AAB99132-AAB99143,
XX AAH75903-AAH75920 and AAG66436.
XX SQ Sequence 822 AA;
alignment_scores:
Quality: 113.00 Length: 475
Ratio: 0.467 Gaps: 20
Percent Similarity: 50.947 Percent Identity: 20.421
alignment_block:
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US-09-303-518D-131 x AAB96063 ..

Align seg 1/1 to: AAB96063 from: 1 to: 822

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238 ProValThrAsnAsnArgAsnGluMetIleAsnAlaSerTrpGlyLe 254
67 .GCCCGCGCATACCGAAGTCCGTTGCGGAGGAATATGTCGGCA 115
|||||:|||||:|||||:|||||:|||||:|||||:|||||:
254 uGlyGluAlaValSerGlyAlaValThrProAspGluTrpIleValG 271
116 TGGCCCGCTCGATGAATAAGAGAGTGAAGCCGTCAAAAGAGCCAA 165
|||: |||: |||: |||: |||: |||: |||: |||: |||: |||:
271 luLysGlyThrTrpLysIleLysGlu...LysValIleAlaLysLysGlu 286
166 GTCTCTTTGACAGACAAAAGATCCGCGCTAGTATTTACTGCGCGGC 215
|||||: |||: |||: |||: |||: |||: |||: |||: |||: |||:
287 ValMet.....ValIleArgAsnProG1 294
216 TTCAGGCAAA.....ATCGCGCTATTACCGTGGC.... 246
|||||: |||: |||: |||: |||: |||: |||: |||: |||: |||:
294 uThrGlyLysGlyThrValThrValLysValAlaGluTrpLeuGlyProG 311
247 .....GAAAGCCGCTACTT.....CAGTCAGTCTGATTGCC 279
|||||: |||: |||: |||: |||: |||: |||: |||: |||: |||:
311 luTrpValGluLysGlnValLeuThrAspGluGlnIleIleGluValAla 327
280 GTTGAAGCAAGACGCAAAATCAGTTCGAACGCTACGTACCTGAAGCGCT 329
|||||: |||: |||: |||: |||: |||: |||: |||: |||: |||:
328 LysMetGlyGlnLysIleGluHisTrpGlyTrpProGlnAspIleG1 344
330 GCAAAATGACAGCGGAAAAAGTGGCGGCAACCTGATTCAATCAGGCT 379
|||||: |||: |||: |||: |||: |||: |||: |||: |||: |||:
344 uTrpAlaTrpAspLysAspGlyLysLeuTrpIleValGlnSerArgp 361
380 TATGACGCTCGCTCGCACCCGCTCGTCAGCAAAATCCCTCGCTAGAT 429
|||||: |||: |||: |||: |||: |||: |||: |||: |||: |||:
361 roValThrThrLeuLysGluThrThrThrGluGluVal... 376
430 GCCGAGCGCTGCGCATCTTCGTCATCGATGGACACCAATCCG...CT 476
|||: |||: |||: |||: |||: |||: |||: |||: |||: |||:
377 ...GluGluAlaGluValIleLeuLysGlyLeuGlyAlaSerProGlyI1 392
477 GCCTCGGACCTACGCTCATCATCAAGAGCCGCGGAGAGCTTCAAAC 526
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392 eGlyAlaGlyArgValValIlePheAspAlaSerGlu..... 405
527 GCGGCTGTGTGATTGAGCGCGCTGACCGAAGCTAAATCCATGCTGTGT 576
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406 .....IleAspLysValLysGlyGluGlyAspValLeuValThr 417
577 AAGCAGCAGCGCGCAGAC...GTGCGCT...GAAAATGCTGCAATAT 620
|||||: |||: |||: |||: |||: |||: |||: |||: |||: |||:
418 ThrMetThrAsnProAspMetValProAlaMetLysArgAlaAlaI1 434
621 CGAACACATGAATTTGCGCGCGCGCTCTGCGCGCTGAGTGGCAGCG 670
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434 elleThrAspGluGlyGlyArgThrSerHisAlaAlaIleValSerArg 451
671 ACATTCATTTTCATCAGCAGCGCGGGAATAAACCGCTGGACCATC 720
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451 luLeuGlyIleProAlaValValGlyThrLysGluAlaThrLysLysLeu 467
721 AATTATCAAGACGATGTCATCGGA..... 747
|||: |||: |||: |||: |||: |||: |||: |||: |||: |||:
468 LysThrGlyAspTrpValThrValAspGlyThrArgGlyLeuValTyrLy 484
748 .....CGTTGTTGTAACAGCGCGCTCTGAATACCGAGCGCGTGG 787
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484 sGlyIleValLysSerLeuValGluLysLysLysLysGluAlaAla 501
788 TTGCTTTGGCGGCGCTGCAAGTCAACAAACGCGCGCTCTTTCGCTACCGTT 837
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501 laAlaProGlyAlaAlaValAlaAlaProLeuValThrGlyLeu 517
838 TTGGTGTGGAAGGTGCTCAACTTACCGCGCGCAATTTGGTTGACGGGA 887
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518 ValLysValAsnValSerMetProGluValAlaGlu..... 529
888 CAACCGCTGATTTCCGGTTCGGTATTGAACGGTSCGATTGCACAAGCG 937
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530 ...ArgAlaAlaAlaThrGlyAlaAspGlyValGlyLeuLeuArgAlaG 545
938 CGCATGATTATTGGCGCGCTACCAATCAGATTCCTGTTATCGAAGA 987
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545 luHisMetIleLeuSerIleGlyGlnHisProValLysPheIleLysGlu 561
988 GCGCGCACAAAGAGCTTTCGGTTCGGCGCGCGGACGCGGACAAATA 1037
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562 GlyLysGluGluLeuValGluLysLeuAlaGluGlyIleGluLysVa 578
1038 CTCATACGCGCGCACCACTCTCGGCCATTT..... 1067
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1068 .CCTAAAAACAACTCTTCAAGTT.....CAGCAGACGCGTCAAC 1107
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594 aProThrAsnGluPheArgGluMetProGlyGlyGluAspGluProGluG 611
1108 GCGCGGACCGCGCGCATGGTACCGATCGGCATTTATGAGCGCGTAATGCC 1157
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611 luArgAsnProMetLeuGlyTrpArgGly.....IleArgArgGly 624
1158 GTTGACATCTCTGCTACCTGCTTTTCGCGATTAACTCTCGCGGATA 1207
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625 LeuAspGlnProGluLeuLeuArg...AlaGluPheLysAlaIleLysLy 640
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640 sValValGluLysGlyTyrAsnAsnIleGly..... 650
1258 GCTTTGTCAGCTCTGCTG.....CCGCGGCAATACGAATACGG 1298
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651 ....ValMetLeuProLeuValSerHisProGluGlnIleArgLysAla 665
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666 LysGluIleAlaArgSerValGly 673
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seq_name: /SIDS1/gcgdata/geneseq/geneseq-emb1/AA2001.DAT:AAB59817

seq_documentation_block:

ID AAB59817 standard; Protein; 999 AA.

XX AAB59817;

XX AC

XX DT

XX 04-APR-2001 (first entry)

XX TutD protein #8.

XX DE

XX KW

XX Toluene degradation; enzyme; waste degradation; TutD.

XX Thauera aromatica.

XX OS

XX OS

XX OS

XX PN

XX XX

XX PD

XX XX

XX PF

XX XX

XX PR

XX XX

(UYOH-) UNIV OHIO.
Coschigano PW;
WPI; 2001-041080/05.
N-PSDB; AAF23625, AAF23627.
Composition comprising toluene degrading enzyme useful for biological treatment of organic compounds, especially for degrading toluene or its analogs -
Disclosure; Fig 5; 122pp; English.
The present invention relates to toluene degrading enzyme genes and proteins tutt (see AAF23629 and AAB59831), tuttI (AAF23630 and AAB59832), tuttF (AAF23631 and AAB59833) and tuttG (AAF23632 and AAB59834). The toluene degrading enzymes are homologues of pyruvate formate lyase. The toluene degrading enzymes are useful for biological treatment of organic compounds and in particular for the degradation of toluene and its analogs contained in liquid or solid waste source. The present sequence is a protein sequence for toluene degrading enzyme, tuttD.
Sequence 999 AA;
SQ

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alignment_scores:
  Quality: 112.50      Length: 536
  Ratio: 0.516        Gaps: 26
  Percent Similarity: 40.672  Percent Identity: 23.694

alignment_block:
US-09-303-518D-131/rev x AAB59817

Align seg 1/1 to: AAB59817 from: 1 to: 999

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1273 .CGAAGCTGCACAAGCAGAGGTCTCTTCGTGCCAATTCACAGCAACCCA.. 1226
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128 pSerArgCysSerSerProSerProArgArgCysProProSerSerProA 145
|||||:|||||:|||||
1225 .....AAGCCTGCGCGGTCTCGGTATCGCGACAGATTAAATCGCGCAAAA 1181
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145 laGlyAlaProGlyAlaThrCysSerArgArgProPheSerArg..... 159
1180 GCAAGGTAGGAGGAGTATCCCAAGGCATTACCGCTCATTAAGTCGCGATC 1131
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160 SerArgaspSerAlaGlyProArgAlaAlaArgPheArgArgCysArgas 176
1130 GGTACCATGGCGGTGCGCGCGGTGTGACGGCTGTCGTCAACTTGAGAG 1081
|||||:|||||:|||||
176 palacysGluArgArgAlaArg..... 183
1080 TTTGTTTTTTAGGAAATGCGCGAGAGTGGTGGCGGTGATGGAGTATTGT 1031
|||||:|||||:|||||
184 .....CysProGlyProArgSerAla 190
1030 CCGGTGCGGCGCAACCCAGCGCAACAGCTCTTTGCTGCGGCTTCTTCG 981
|||||:|||||:|||||
191 ProSerIleArgArgGlySerArgaspArgSerArgAlaSerArgSerAr 207
980 ATAACGGAAATCTGAT...TGTTGTAGCGTCCCAATAATCATCGCGGCC 934
|||||:|||||:|||||
207 gSerArgGlySerProLeuCysGlyAlaThrAlaThr..... 219
933 TTGTGCAATCGCACCGTTCAATACCGAAGCGGAAA..... 899
|||||:|||||:|||||
220 .....SerCysProArgArgArgArgCysSerIleGly 230
988 ...TCACGGCGGTGTCCGCGTCAACCAATTCGCGCGGTGAAGTGGAGC 852
|||||:|||||:|||||

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157 .....TTTTACGGCTTCACCTTCCTTGATTTT 130
158 .....| | | | |
516 oSerSerProLeuAsnArgProPheAlaArgSerAlaPro.....A 531
129 CATCGAGGGCGCATCCGACATATCTTCGCCAAGCAAGCGGACTCGG 80
130 :| | | | |
531 laSerThrProCysArgArgHisAsnArgArgTyrGlySerArgArg 547
79 .....TAATGGCGGGCGCGTCATAAA..... 59
548 ProPheArgArgPheAlaCysSerTrpSerSerGlnHisAspProAl 564
58 .....TGACTTGTCCGGTCTGCCCGC 37
564 aSerGlnAspProGlnArgGlyThrCysProLeuArgAsnAlaCysProG 581
36 GATGGGCA 29
581 lyTrpAla 583
seq_name: /SIDS1/gcgdata/geneseq/geneseq-emb1/AA1999.DAT:AAV28654
seq_documentation_block:
ID_ AAV28654 standard; Protein; 802 AA.
XX
AC AAV28654;
XX
01-OCT-1999 (first entry)
XX
DE Murine Cytoplasmic phosphatase, 270PEP protein.
XX
KW Lymphoid protein Tyrosine Phosphatase; Lyp protein; lymphoid cell;
KW intracellular tyrosine phosphatase; ptpase; lymphocyte; murine;
KW protein tyrosine kinase; ptk; immunosuppressant; pEST sequence;
KW T cell antigen receptor signalling; autoimmune disease; transplant;
KW cytokine receptor signalling.
XX
OS Mus sp.
XX
FH Key
FH Domain 27..288
FH /label= ptpase_domain
FH /note= "Catalytic protein tyrosine phosphatase domain"
FH Binding-site 613..621
FH /label= SH3_binding_site
FH /note= "Proline rich sequence"
FH Binding-site 689..695
FH /label= SH3_binding_site
FH /note= "Proline rich sequence"
FH Binding-site 790..798
FH /label= SH3_binding_site
FH /note= "Proline rich sequence"
XX WO936548-A1.
XX
XX 22-JUL-1999.
XX
XX 18-JAN-1999; 99WO-CA00038.
XX
XX 16-JAN-1998; 98CA-2220853.
XX (HSCR-) HSC RES & DEV LP.
XX
XX Roifman CM;
XX
XX WPI; 1999-444404/37.
XX
XX New nucleic acid encoding intracellular tyrosine phosphatase and
XX related proteins, used to modulate signalling through T cells,
XX particularly as immunosuppressant
XX
XX Disclosure; Page 63-64; 105pp; English.
PS

```

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XX
CC The present protein sequence is that of the murine phosphatase, 270PEP
CC that has a single catalytic domain. The non-catalytic portion of the
CC phosphatase contains unique sequences, including five PEST sequences
CC rich in Pro, Glu or Asp, Ser and Thr. 270PEP shares about 70% sequence
CC identity with the human cytoplasmic phosphatase LypL. Lyp proteins are
CC important for regulation of T cell antigen and cytokine receptor
CC signalling and for early and late stages of T cell differentiation.
CC 270PEP has immunosuppressive activity. Compounds that increase
CC expression of Lyp protein can be used as immunosuppressive agents to
CC reduce or prevent T cell activation or proliferation, to control
CC thymocyte differentiation, to treat autoimmune diseases and transplant
CC situations.
XX
SQ Sequence 802 AA;

```

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alignment_scores:
  Quality: 112.00      Length: 351
  Ratio: 0.622        Gaps: 20
Percent Similarity: 51.282 Percent Identity: 24.501

alignment_block:
US-09-303-518D-131/rev x AAV28654 ..
Align seg 1/1 to: AAV28654 from: 1 to: 802

```

```

955 GTCCCAATATCAT...GCGCGCTTGTGCATTCGCACCGTTCATATACC 909
956 :| | | | |
957 lSeSerAspAsnHisLeuGlyArgGluIleGlnAlaGlnCysSerIlePr 313
958 :| | | | |
959 :| | | | |
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961 :| | | | |
962 :| | | | |
963 :| | | | |
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999 :| | | | |
1000 :| | | | |

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```
1009 CGAACAGCTCTTTGCTGCGCCCTTCTTCGATACGAGGAAATCTGATTGTGG 960
||||| :|: |||||:|:|
210 roAsn...LysCysTyrAlaProSerAlaIlePro..... 220
959 TAGCGTCCCAATATATCATCGCGCTTGTGCAATCGCACCGCTTCAATAC 910
||||| :|: |||||:|:|
221 ...ThrProGlnArgThrSerThrPro...GlyLeuAlaLeuPheProG1 235
909 CGAACCGGAAATCAGCGGTGTCGCGCPCAAAC..... 876
||||| :|: |||||:|:|
235 yProProSerProValAlaAsnSerThrProLeuThrLeuProVal 252
875 ..AATTGCGCG.....GCGTAAGTTGAGACACCTTCGCACCCAAA 837
||||| :|: |||||:|:|
252 alGlnSerProLeuAlaThrAlaAlaSerAlaSerThrSerAlaProVal 268
836 ACGGTACGCAAGAGCGCGTGTGTGACTTCGACGCGCCGCCAAGCAAC 787
||||| :|: |||||:|:|
269 SerCysGlySerSerAlaSerLeuLeuArgGlyProHisProGlyThrSe 285
786 CACGCGCTCGGTATTACAGCGCGCTTACGACAAACGTCGATAGCAA 737
||||| :|: |||||:|:|
285 rAspLeuHisIleSerThrProAlaAlaThrThrLeuProVal..... 300
736 TCACGTCTGTGATTAATGATGTCACACGCTTTATTCGCGCCGACTGGC 687
||||| :|: |||||:|:|
301 .....MetIleLysThrGluProThrSerProThrPro 311
686 TCGATGAATGAATGTCGCTGCCACTCAAG...CGGCGAGGATGGGGCC 640
||||| :|: |||||:|:|
312 Ser.....AlaPheLysGlyProSerHisSerGlyAs 322
639 GCCAATTCATGTGTTTCGATATTCGACGATTTTCAGACGCGAGCTGTG 590
||||| :|: |||||:|:|
322 nProSerHisGlyThrLeuGlyLeuSer.....GlyThrLeuG 335
589 CGCTGCTGCTTACACACATGGAATTTACGTTGCGTCAGCGCTCAAT 540
||||| :|: |||||:|:|
335 lYArgAlaTyrThrSerThr..... 341
539 ACCAACAGCGCGCTTTGAAGTCTTCGCGGCTTCTTTGATGATGACGGT 490
||||| :|: |||||:|:|
342 .....SerValProIleSerLeuSerAlaCysLe 351
489 AGGTGCGCAGCGGATGTTGTCATCGCATTCAGCAAGATGGCGA 440
||||| :|: |||||:|:|
351 uAsnProAlaLeuSerGlyLeuSerSerSerThrPro.....LeuA 366
439 ACGCTCGGCATCTACGCGAGGATTTGCTGAACGCGGTCGCGAAGC 390
||||| :|: |||||:|:|
366 sNGlySerAsnProLeuSerSerIleSerLeuProProHisGlySerSer 382
389 CAGTCCATAAGCTGATTAATCAGTTGCGGCGCATTTCGCTGCT 340
||||| :|: |||||:|:|
383 ThrProIleAlaProValPheThrAlaLeuProSerPheThrSerLeuTh 399
339 CAATTTGCCAGCGCTTACGATAGCTGCGTTCGAACCTCGATTTCGCTG 290
||||| :|: |||||:|:|
399 rAsnAsnPheProLeuThrGlyAsnProSerLeuAsnProSerValSerL 416
289 TGCCTTCAACGCAATCAGCACTACGATGAGTACGCGCTTTTCG..... 246
||||| :|: |||||:|:|
416 euProGlySerLeuIleAlaThrSerThrAlaAlaThrSerThrSer 432
245 .....CCACGGTGAATAGCGGCTTTTG..... 222
433 LeuProHisProSerThrAlaAlaValLeuSerGlyLeuSerAlaSe 449
221 ....CCTGAACCGCGCA..... 207
449 rAlaProValSerAlaAlaProPheProLeuAsnLeuSerThrAlaValP 466
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seq_name: /SIDS1/gcgdata/geneseq/geneseq-emb1/AA1999.DAT:AA1999

seq_documentation_block:

ID AAY31745 standard; Protein; 430 AA.

XX AAY31745;

XX 22-NOV-1999 (first entry)

XX Mycobacterium tuberculosis specific DNA-encoded polypeptide.

XX Tuberculosis; infection; diagnosis; DNA probe.

XX Mycobacterium tuberculosis.

Key Location/Qualifiers

FT Misc-difference 4 /note= "encoded by TGA"

FT Misc-difference 6 /note= "encoded by TGA"

FT Misc-difference 20 /note= "encoded by TGA"

FT Misc-difference 29 /note= "encoded by TGA"

FT Misc-difference 54 /note= "encoded by TGA"

FT Misc-difference 64 /note= "encoded by TGA"

FT Misc-difference 69 /note= "encoded by TGA"

FT Misc-difference 89 /note= "encoded by TGA"

FT Misc-difference 99 /note= "encoded by TGA"

FT Misc-difference 114 /note= "encoded by TGA"

FT Misc-difference 119 /note= "encoded by TGA"

FT Misc-difference 129 /note= "encoded by TGA"

FT Misc-difference 159 /note= "encoded by TGA"

FT Misc-difference 169 /note= "encoded by TGA"

FT Misc-difference 182 /note= "encoded by TAG"

FT Misc-difference 185 /note= "encoded by TGA"

FT Misc-difference 219 /note= "encoded by TGA"

FT Misc-difference 259 /note= "encoded by TGA"

FT Misc-difference 269 /note= "encoded by TGA"

FT Misc-difference 291 /note= "encoded by TGA"

FT Misc-difference 323


```

367 yArgAlaProAlaCys..... 372
633 ATTTGGCGCGCCGATCTCGCGGCTTGAGTGGCAGCAGACATTCA 682
373 .....CysAsp 374
583 TCGAGCCAGTCGGCGCAATAAACCCTGTGGACCATCAATTATCAAGAC 732
375 ArgValGlnThrPro***HisArgPro..... 383
733 GTGATTGTATCGGACGCTTTGTCGTAAACAGCGCTCTGAATACCGAGCG 782
384 .....ProIleSerValProThrS 390
783 CGTGGTTG...CCTTTGGCGGCGTCGAAGTCAACAACCGCGCTCTTGC 829
390 erTrpCysGlyProTrp***Thr...ProAlaProArgThr***TrpCys 405
830 GTACCG.....TTTGGTGCAGAGGTGCTCAACTTACCGCGCGGAA 873
406 CysProMetAlaMetTrp.....ProProLysAs 415
874 TTGG.....TTGACGCGGACAAACCGCGTGTATTCGGTTCGGTATT 914
415 nTrpTrpProGlyValProArgArgSerAlaGlyAlaSerThrTrpTyr. 431
915 GAACGGTGCATGTCACAAAGCGCGCATGATTATTGGGACGCTACCACA 964
432 .....ProCysArgProAspArgTrpCysArgGlyTrp 442
965 ATCAGATTTCGTTATCGAAGAGCGCGCAGCAAGAGCTTCGGCT... 1012
443 ProArgTrpProCysMetThrArgProAlaArgProSerThrThrAlaTh 459
1013 .....GGGTTGCGCGC..... 1024
459 rAlaTrpProValProValLeuProGlyThrAspArgCysAlaLeuP 476
1025 ..AGCGGACAAATCTCCATCAGCGCGCACCCTCTCGGCC..... 1063
476 roProLysArgArg***ProGlyProValProAlaSerArgAlaThrVal 492
1064 ...ATTTCCTAAAAACAACTCTTCAAGTTCA.....CG 1095
493 TrpValSerArgAlaThrArgCys***SerSerProThrMetSerProAr 509
1096 ACAGCGCTCAACGGCGGCGACCGGCCCATGTGTACCGATCGGCACCTATGA 1145
509 gArgProSerValTrpSerThrCysCysTrpHisArgGluAlaIle.... 524
1146 GCGCGTAATGCGGTGGACATCTCTACCTGCTTTTGGCGGATTTAA 1195
525 .....Trp***ArgCys..... 528
1196 TCGTCGGCGATACCGACAGCGCGAGCTTTGGTT...GCTTGAATTG 1242
529 ...***LeuAlaProAla***ProLysThrTrpLeuSerSerTrpAsnG1 544
1243 GACGAAGACCTCGCTTTGTGGAGCTTCTCTCCCGCGGCAATACGA 1292
544 yMetCysThrThrThrIleGlnAlaProSerTrpSerProThrAlaProA 561
1293 ATACGCGC.....CGCTGT 1306
561 spThrAlaAlaThrArgCys 567
seq_name: /SIDSI/gcgdata/geneseq/geneseq-emb1/AA2001.DAT:ABB64198
seq_documentation_block:
ID: ABB64198 standard; Protein: 2406 AA.
XX
AC ABB64198;
```

```

XX
DT
XX
DE Drosophila melanogaster polypeptide SEQ ID NO 19386.
XX
KW Drosophila; developmental biology; cell signalling; insecticide;
pharmaceutical.
XX
OS Drosophila melanogaster.
XX
PN WO200171042-A2.
XX
PD 27-SEP-2001.
XX
PF 23-MAR-2001; 2001WO-US09231.
XX
PR 23-MAR-2000; 2000US-191637P.
PR 11-JUL-2000; 2000US-0614150.
XX
PA (PEKE ) PE CORP NY.
XX
PI Venter JC, Adams M, Li PWD, Myers EW;
XX
WPI; 2001-656860/75.
DR N-PSDB; ABL08301.
XX
PT New isolated nucleic acid detection reagent for detecting 1000 or more
genes from Drosophila and for elucidating cell signalling and cell-cell
interactions
XX
PS Disclosure; SEQ ID NO 19386; 2lpp + Sequence Listing; English.
XX
CC The invention relates to an isolated nucleic acid detection reagent
capable of detecting 1000 or more genes from Drosophila. The invention is
useful in developmental biology and in elucidating cell signalling and
cell-cell interactions in higher eukaryotes for the development of
insecticides, therapeutics and pharmaceutical drugs. The invention
discloses genomic DNA sequences (AB1616176-AB1616175), expressed DNA
sequences (AB1616176-AB1616175) and the encoded proteins
(ABB57737-ABB72072).
XX
CC The sequence data for this patent did not form part of the printed
specification, but was obtained in electronic format directly from WIPO
at ftp.wipo.int/pub/published_pct_sequences.
XX
SQ Sequence 2406 AA;

alignment_scores:
Quality: 108.00 Length: 473
Ratio: 0.519 Gaps: 28
Percent Similarity: 43.975 Percent Identity: 22.833

alignment_block:
US-09-303-518D-131/rev x ABB64198 ..
Align seg 1/1 to: ABB64198 from: 1 to: 2406

1298 CCGTAT.....TCGTATTCGCCGGCAGACGAGCT 1267
|||||
1266 ProTyrSerGlyLeuThrHisGlySerTyrLeuProProValLeuProVa 1282
1266 G.....CACAAAGCGAGGT 1253
1282 lAlaThrProAsnLeuSerAsnLeuProProThrGlnHisArgSer...S 1298
1252 CTTCTTCGTCCTCAATTCACAGCAACCCAAAGCTCGCGCTGTCGGTATCG 1203
|||||
1298 erAspSerArgAsnSerArgGluSerProAla..... 1308
1202 CCGACGATTAAATCGCGCAAAAGCAAGGTAGC....AGGATGTCCACGG 1156
|||||
1309 ...SerLeuLysSerThrProSerAsnIleGlyLeuAsnValSerMetAl 1324
```


CC is a protein sequence encoded by toluene degrading enzyme gene, TtutD/E.

XX
SQ Sequence 1592 AA;

alignment_scores:

Quality: 109.50 Length: 550
Ratio: 0.517 Gaps: 26
Percent Similarity: 38.545 Percent Identity: 22.727

alignment_block:

US-09-303-518D-131/rev x AAB59827 ..

Align seg 1/1 to: AAB59827 from: 1 to: 1592

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1312 TGCACACAGCGCGCCGATTCGATTTGCGCGGAGCA..... 1274
|||||:||||| ||| ||| |||
707 CysThrSerArgGlyArgSerArgCysSerProAspArgCysCysArgTr 723
1273 CGAAGCTGCACAAAGCGAGGTCTTCGTCCTCAATCCAAAGCAACCCA. 1226
|||||:||||| ||| ||| |||
723 pSerArgCysSerSerProSerProArgArgCysProProSerSerProA 740
1225 .....AAGCGTCGCGCTGTCGCTATCGCGAGCATTAATCGCGCAAAA 1181
|||||:||||| ||| ||| |||
740 LaGlyAlaProGlyAlaThrCysSerArgArgProPheSerArg..... 754
1180 GCAAGGTAGCGAGGATCTCAACGGCANTACGGCTCATAGTGGCGATC 1131
|||||:||||| ||| ||| ||| ||| ||| |||
755 SerArgAspSerAlaGlyProArgAlaAlaArgPheArgArgCysArgas. 771
1130 GTTACCATGCGCGCTGCGCGCGCTTGCACGCTGCTGTAACCTGAAGAG 1081
|||||:||||| ||| ||| |||
771 PaLysGluArgAlaArg..... 778
1080 TTTGTTTTTGGAAATAGGCGAGAGTGGTGGCGGTGATGAGTATTTGT 1031
|||||:||||| ||| ||| |||
779 .....Cys 779
1030 CCG.....GC 1026
780 ProGlyProArgSerAlaProSerIleArgArgGlySerArgAspArgSe 796
1025 TCGCGCGCAACCCAGCGAGAGCTCTTTGCTGGCGCTTCTTCGATAAC 976
|||||:||||| ||| ||| ||| |||
796 rArgAlaSerArgSerArgGlySerProLeuCysGly..... 808
975 GGAATCTGATGTGGTAGCTGCCAATAATCATGCGCGCTTGTGCAA 926
809 .....Ala 809
925 TCGCACCGTTCAATACCGAACCGGAAA.....TCACG 894
|||||:||||| ||| ||| ||| |||
810 ThrAlaThrSerCysProArgArgArgArgCysSerIleGlyAlaSerSe 826
893 CGGTTGTCGGCTCAACCAATTCGCCCGGTAAGTTGAGACACCTTCGC 844
|||||:||||| ||| ||| ||| ||| |||
826 rGlyCysProHisProProValArgArgSerProValAsnSerSerIysA 843
843 ACCCAAAACGGTACGCA.....AGAGCGCGGTT 815
|||||:||||| ||| ||| ||| |||
843 rGAlaHisArgArgCysThrAlaArgArgGlyArgPheArgGlyProThr 859
814 TGTTCACCTGACGGCGC.....CCAGGCAACACAC 783
|||||:||||| ||| ||| ||| |||
860 SerArgAspThrGlyArgArgCysTrpArgTrpProArgProArgAr 876
782 CGCTCGG.....TATTCAGCGCGCTGT 760
|||||:||||| ||| ||| ||| |||
876 gCysArgCysSerArgArgTrpGlyArgProLeuTrpAlaSerGlyCysp 893
759 TACGAACAACCTCGATAGCAATCACTCTTGATAATTGATGTCACACA 710
|||||:||||| ||| ||| ||| ||| ||| |||
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893 roArgAlaArgTrpArgArgGlySerAsn.....TipSerSer 905
709 .....CGGTTTTTATTCGCGCGCGACTG... 689
906 GlyArgSerSerAlaAlaSerProLysArgThrCysGlyArgArgValar 922
688 .....GCTCGATGAATGTGCGTGCACCTCAAG..... 657
922 gSerAspThrSerAlaArgSerArgCysProAlaSerSerProIlea 939
656 .....CCGGCAGGATCGGCGCG 639
939 rgTrpThrGlyArgCysArgArgTrpArgProLeuGlyCysSerPro 955
638 CCAAAATTCATGTTCGATATTGCGACATTTTCAGACAGCGCAGCTGTC 589
956 ArgAlaThrCysThrAla.....ArgCysGlyArgAspGlyCysSerAl 970
588 G.....CCTGCTGCTTTACACACATGATTTTACGTTTCGTCAGCGCGC 545
970 aPhePheGlyAsnProLeuHis.....ArgSerLeuArg.... 981
544 TCAATACCAACAGCGCGCTTGAAGTCTTCGGCGCTTCTTTGATGATG 495
981 ..... 981
494 ACCGTAGGTGCGCAGCCAGCGG ATTGGTGTCATCGCATTTGACGAAGA 446
982 .....GlyProTrpAlaAlaProPheArgAlaHisArgSerArgSerTh 996
445 TGGCGAACCGCTCGCATCTACGCGAGGATTTCTGTAACGCGCGGTG 396
996 rThrArgArgCysAlaValArgGlySerSerArgHisAspArgThrAlas 1013
395 CGAAGCGCAGTCCATAAGCTGATTGAATCAGTTTTCGCGCGCATTTC 346
1013 erThrArgArgProHisLysProProLysGlyCysAlaThrAspIleHis 1029
345 GCTGCTCAATTTTGCCA.....GCGCTTCA 321
1030 SerGlyArgTy-CysTrpProArgThrAlaSerSerArgAlaAlaSerGl 1046
320 GGTACGTACGCTCGNACTCGATTTCGCGT.....TGCCTTCAACGCG 277
1046 yAlaSerAlaLysArgThrArgLeuArgArgArgSerCysProValArgS 1063
276 AATCAGCACTGACTGAAGTACGCGCTTTTCGCCAC..... 242
1063 exProArgArgArgGlyThrArgAlaAlaTrpHisSerAlaCysGlySer 1079
241 GGTGAATAGCGGATTTTTCGTCGAGCGCGCAGTAATACTACGCGCC 192
1080 SerSerArgArgProSerSerGlyArgProTrpSerValProIleArgPr 1096
191 GGATTCCTTTTCTTCAACAGCACTTGCCTT..... 158
1096 oSerSerIleCys...GlyArgAlaValGlyLeuThrSerProSerSerP 1112
157 .....TTTTCACGCGCTTTCACCTTCCTTGTGATTTTCATCGAGG 122
1112 roLeuAsnArgProPheAlaArgArgSerAlaPro.....AlaSerThr 1126
121 GGGCGATGCGCAGATATTCTTCGCAAGCAACGCGACTTCGG..... 80
1127 ProCysArgArgHisAsnArgArgArgTyGlySerArgArgProPheAr 1143
79 .....TAATGGCGCGCGCTCATAAA..... 59
1143 gArgArgPheAlaCysSerTrpSerSerGlnHisAspProAlaSerGlnA 1160
58 .....TGACTTCTCGCGCTGTCGCGCGAGTGGCA 29
1160 spProGlnArgGlyThrCysProLeuArgAsnAlaCysProGlyTrpAla 1176
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seq_name: /SIDSL/gcdata/geneseq/geneseq-emb1/AA2001.DAT: AAB27242
seq_documentation_block:
ID   AAB27242 standard; Protein; 571 AA.
XX  AC   AAB27242;
XX  DT   27-MAR-2001 (first entry)
XX  DE   Human EXMAD-20 SEQ ID NO: 20.
XX  KW   Extracellular matrix and adhesion-associated protein; EXMAD; cancer;
XX  KW   inflammation; reproductive disorder; cardiovascular disorder;
XX  KW   immune disorder; musculoskeletal disorder; developmental disorder;
XX  KW   gastrointestinal disorder; cell proliferation disorder.
XX  OS   Homo sapiens.
XX  XX   WO200068380-A2.
XX  PD   16-NOV-2000.
XX  PF   10-MAY-2000; 2000WO-US12811.
XX  PR   11-MAY-1999; 99US-0133643.
XX  PR   23-AUG-1999; 99US-0150409.
XX  XX   (INCY-) INCYTE GENOMICS INC.
XX  XX   Bandman O, Hillman JL, Tang YT, Lal P, Yue H, Baughn MR, Lu DAM;
XX  XX   Azimzai Y;
XX  DR   WPI; 2001-007395/01.
XX  DR   N-PSDB; AAC66909.
XX  PT   Isolated polynucleotide encoding extracellular matrix or
XX  PT   adhesion-associated protein (EXMAD) useful for diagnosing, treating, or
XX  PT   preventing disorders associated with expression of EXMAD such as
XX  PT   proliferative, immune and genetic disorders -
XX  XX   Claim 1; Page 106-107; 129pp; English.
XX  CC   The present invention provides the protein and coding sequences for 25
XX  CC   novel extracellular matrix and adhesion-associated proteins (EXMADs).
XX  CC   These are designated EXMAD-1, EXMAD-2, EXMAD-3, EXMAD-4, EXMAD-5,
XX  CC   EXMAD-6, EXMAD-7, EXMAD-8, EXMAD-9, EXMAD-10, EXMAD-11, EXMAD-12,
XX  CC   EXMAD-13, EXMAD-14, EXMAD-15, EXMAD-16, EXMAD-17, EXMAD-18, EXMAD-19,
XX  CC   EXMAD-20, EXMAD-21, EXMAD-22, EXMAD-23, EXMAD-24 and EXMAD-25. They are
XX  CC   useful in the prevention and treatment of cancers, cell proliferation,
XX  CC   cardiovascular, reproductive, immune, musculoskeletal, developmental and
XX  CC   gastrointestinal disorders and inflammation.
XX  SQ   Sequence 571 AA;

alignment_scores:
Quality: 106.50      Length: 514
Ratio: 0.467         Gaps: 19
Percent Similarity: 44.358      Percent Identity: 18.872

alignment_block:
US-09-303-518D-131/rev x AAB27242
Align seg 1/1 to: AAB27242 from: 1 to: 571

1340 CCTCTCTCTCAATGTTTCAGCAGCTTTCGCAACAGCGGCGGATTC 1291
||| ||| :|||:||||| :|||:||||| :|||:||||| :|||:|||||
23 ProMetMetProThrThrSerGlyThrSerGlnAlaSerSerPheAs 39
1290 GTATTTCCCGGACAGAGCTGACAAAGCGAGGTCTCTCTCTGTC 1241
: :|||:||||| :|||:||||| :|||:||||| :|||:|||||
39 nThrAlaLysThrSerThrSerLeuHis.....SerHisThrSer 53
```

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1240 ATTCACAGCAACCAACCCCTGCGCGTGTGCGTA..... 1206
erThrHisHisProGluValThrProThrSerIleThrAsnIleThrLeu 69
1205 .....TCGCCGACGATTAAATCGCGCAAAAG 1180
70 AsnProThrSerIleGlyThrThrProValAlaHisThrThrSerAl 86
1179 CAAGTAGGAGGATGCCAACGGCATACGCGCTCATAGTCCGCGATC 1130
: :|||:||||| :|||:||||| :|||:||||| :|||:|||||
86 aThrSerSerArgLeuThrThrProPheThrThrHisSerProThrG 103
1129 GTACC.....ATGGCGCGTCCCGCCGCTGACGCTGTCGTAACCTG 1086
|||||:||||| :|||:||||| :|||:||||| :|||:|||||
103 lYserSerProIleSerSerThrGlyProMetThrAlaThrSerPheGln 119
1085 AAGAGTTTGTTTTATAGAAATGCCGAGAGTGGTGGCGTGATGGAGTA 1036
: :|||:||||| :|||:||||| :|||:||||| :|||:|||||
120 ThrThrThrTyrThrPro..... 126
1035 TTGTGCGGCTGCGGCGCAACCCAGCGAACAGCTCTTTGTCGCGCCTT 986
: :|||:||||| :|||:||||| :|||:||||| :|||:|||||
127 .....ProSerHisProGlnThrThrLeu..... 134
985 CTTTCGATAACGGAATCTGTATTCTGTAGGTCCCAATAATCATCGCG 936
134 ..... 134
935 CCTGTGCAATCGACCGCTTCAATACCGAA.....CCGGAATC... 897
||| :|||:||||| :|||:||||| :|||:||||| :|||:|||||
135 ProThrHisValProPheSerThrSerLeuValThrProSerThrHi 151
896 .....ACGCGGTTGTCGCGCTCAACCAATTCG 869
|||:||||| :|||:||||| :|||:||||| :|||:|||||
151 sThrValIleThrThrHisThrGlnMetAlaThrSerAlaSerIleH 168
868 CGCGCGTAACTTCAGACACTTCGCGCACCAACAGCTACGAGAGCGCG 819
|||:||||| :|||:||||| :|||:||||| :|||:|||||
168 isSerThrProThrGlyThrValProProThrThrLeuLysAlaThr 184
818 GGTTCGTTGCTGCGCGCGCAACGCGCTCGGTATTCAG 769
|||:||||| :|||:||||| :|||:||||| :|||:|||||
185 GlySerThrHisThrAlaProMetThrValThr..... 196
768 ACGCGCTGTACGAACAAACGTCGCGTACATCAGCTTTCATATGA 719
|||:||||| :|||:||||| :|||:||||| :|||:|||||
197 .....ThrSerGlyThrSerG 202
718 TGTTCACACGCGTTTATTCGCGCGACTGCGTCAATGAAATGAATGTC 669
|||||:||||| :|||:||||| :|||:||||| :|||:|||||
202 lNThrHisSerSerPheSerThrAlaThrAlaSerSerPheIleSer 218
668 .....GTGCCACTCAAGCGCGAGGATCGGCGC 640
: :|||:||||| :|||:||||| :|||:||||| :|||:|||||
219 SerSerSerTrpSerSerTrpLeuPro...GlnAsnSerSerSerArgPr 234
639 GCCAATTCATGTGTTTCG..... 621
|||||:||||| :|||:||||| :|||:||||| :|||:|||||
234 OProSerSerProIleThrThrGlnLeuProHisLeuSerSerAlaThr 251
620 .....ATATTGGCAGCATTTTTCAGACGGCAGC 594
: :|||:||||| :|||:||||| :|||:||||| :|||:|||||
251 hrProValSerThrThrAsnGlnLeuSerSerSerPheSerProSerPro 267
593 TCTGCGCGCTGCTGCTTACACACATGGATTTTACCTTCGTCAGCGGCT 544
|||||:||||| :|||:||||| :|||:||||| :|||:|||||
268 SerAlaProSerThrValSerSerTyrValProSerSerHisSerSerPr 284
543 CAATACCAACAGCGCGCTTGAAGTCTTCGCGCGCTTCTTTGATGATGA 494
: :|||:||||| :|||:||||| :|||:||||| :|||:|||||
284 oGlnThrSerProSerValGlyThrSerSerSerPheValSerAlaP 301
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CC diagnostic amino acid sequences of the invention.
CC Note: the sequence data for this patent did not appear in the printed
CC specification, but was obtained in electronic format directly from WIPO
CC at ftp.wipo.int/pub/published_pct_sequences.
XX
SQ sequence 1078 AA;

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alignment_scores:
  Quality: 106.50      Length: 486
  Ratio: 0.461        Gaps: 21
  Percent Similarity: 47.531  Percent Identity: 21.811

alignment_block:
US-09-303-518d-131/rev x ABG21954
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320 AlapheAlaLeuValThrSerSerLeuSerGluSerValPheLeuArgAs 336
1301 .....GGGCGGATTCGATTTCGCCGGGAGAGCAAGCTGCACAAAG 1259
:::|:::|:::|:::|:::|:::|:::|:::|:::|:::|:::|
336 pValPheGlyLeuSerAlaPheLeu.....ArgSerLeuAlaLysS 350
1258 CGAGGCTCTTCTCGTCCAAATTCACCAAGCAACCCAAAGCTCGCGCTGTCG 1209
:::|:::|:::|:::|:::|:::|:::|:::|:::|:::|:::|
350 eAsnSerLeuGlyLysIleSerPheLeuSerSerSerLeuLeuAlaSer 366
1208 GTATCGCGCGAGTAAATCG.....CGCAAAAG 1180
:::|:::|:::|:::|:::|:::|:::|:::|:::|:::|:::|
367 AlaserValThrIleSerSerProSerThrSerSerLeuGlnArgLysAl 383
1179 CAAGGTAGGAGGATG...TCCACGCGCATACCGCTCATAGTGCAGA 1133
:::|:::|:::|:::|:::|:::|:::|:::|:::|:::|:::|
383 aSerPheAlaArgLeuProSerSerGlyLeu.....ProS 395
1132 TCGGTACCATCGCGCGCGCGCTGAGCGTGTGCGTGAACCTGAAG 1083
:::|:::|:::|:::|:::|:::|:::|:::|:::|:::|:::|
395 eGluThrCysIleArgLeuAlaSerAlaSerProValLysGlyLeuLys 411
1082 AGTTTGTTTTAGGAATGCCGAGAGTGGTGGCGCTGATGGAGTATTT 1033
:::|:::|:::|:::|:::|:::|:::|:::|:::|:::|:::|
412 ArgVal.....SerValLeuIleThrSerPheLe 421
1032 GTCCGGCTGCGCGCAACCCAGCGACAGCTCTTTGCTCGCGCTCTTT 983
:::|:::|:::|:::|:::|:::|:::|:::|:::|:::|:::|
421 uSerLeuAlaGlySerSerProProArgIleAlaSerSerGluProSerS 438
982 CGATAACGGAATCTGATTGGTGGTAGCTCCCAATAATCATGCGCGCT 933
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438 eLeuPheASP.....AlaVal 443
932 TGTGCAATCGCACCGTTCAATACCGAACCGGAATCATCGGGTTGTCGC 883
:::|:::|:::|:::|:::|:::|:::|:::|:::|:::|:::|
444 SerSerMetGlyProValSerSerArg.....SerMetAspAl 456
882 GTCAACCAATTCGCGGGGGTAAAGTTGAGACACCTTCGCCACCCAAACGG 833
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456 aSerSerSerLeuSerProThrSer..... 464
832 TAGCAAGAGCGCGGTTGTGACTTGCAGGCGCGCCCAAGCAACCAACG 783
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465 .....LeuThrAlaThrProProSerAlaSerPro 474
782 CGCTCG.....GTATTGAGCGGCTGTTTACGAACAACCTCCGAT 742
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475 SerSerAspValAlaAlaSerLeuArgAlaAlaThrSer***ArgCysSe 491
741 ACCAATCAGCTTCTGATTAATGATGGTCCACACGGTTTATTTCGCGCGA 692
:::|:::|:::|:::|:::|:::|:::|:::|:::|:::|:::|
491 rLysPheArgAla.....AlaAlaSerPheAlaSerT 502
```

```
691 CTGGCTCGATGAATGATGTGCGTGCACCTCAAGCGCGCAGGATCGGG 642
:::|:::|:::|:::|:::|:::|:::|:::|:::|:::|:::|
502 hAlaPheSerLys.....ProCysSerProSerGluAlaAla 514
:::|:::|:::|:::|:::|:::|:::|:::|:::|:::|:::|
641 ...CGGCCAAATTCATGTGTTTCGATATTCGACAGATTTTCAGACGGCAC 595
:::|:::|:::|:::|:::|:::|:::|:::|:::|:::|:::|
515 LeuSerValSerSerThrValSerIleLysAlaCysPheSerGlyAlaSe 531
594 GTCTCGCGCTGCTGCTTTACACATGATTTTACGTTTCGTCAGCGCGC 545
:::|:::|:::|:::|:::|:::|:::|:::|:::|:::|:::|
531 rSer.....LeuLeuArgProThrAlaSerP 540
544 TCAATACCAACAGCGCGGTTTGAAGTCTTCGCGCGCTTCTTTGATGATG 495
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540 heAlaAlaAlaCysProMetGlySerAspProProSerProSerLeuLeu 556
494 ACCGTAGGTGCGGACGCCAGC.....GG 472
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557 ThrProSerAlaSerAlaAlaProSerAlaProLeuSerMetProGlygl 573
471 ATTGTGTCCATCGCATTCGACGAGATGCGAAGCGCTCGGCATCTACGG 422
:::|:::|:::|:::|:::|:::|:::|:::|:::|:::|:::|
573 yIleSerLeuPheSerValAsnSerSerAlaAsnSerGlyCysSerAlav 590
421 CAGGGATT...TTGTGAACGAGCGGGTGCAGGCGCAGTCCATTAAGCCT 375
:::|:::|:::|:::|:::|:::|:::|:::|:::|:::|:::|
590 alGlyPheHisAlaLeuLysGlyAlaSer***ThrAlaProAlaSerPro 606
374 GATTGAATCAGGTTCGGCGGCACCTTTTCGCTGCTCAATTTTCCGACGGC 325
607 His.....SerAlaAlaSerProPheProCysThrMetSerAlaSerLe 621
324 TTCAGGTAGCTAGCGTTCGACATTCGATTTCGCTTGCCTTCAACGGCAA 275
621 ucysrtrpGlyAlaSerSerSerIleAlaSerPhePheSerSerValt 638
274 TCACGACTGACTGAAGTACGCGCTTTTCGCCACGGTGAATACCGCGGATT 225
638 yrVal.....PheCysThrAsnCysTrpAspSerLeu 648
224 TTCCTGTAAGCGCGCGCTAATACTACGCCCGGATCTTTTTCG..... 180
649 PheGluLeuIleAlaAlaValSerSerThrMetLysPheIleValGlyAl 665
179 .....TCTTCAACAGCACCTTGGCTTTTTCACGGCTT 146
665 aProArgAlaArgThr***AlaAsnGlyAsnGluSerPheLeuProLeuV 682
145 CACCTTCTCTGATTTCATCGAGGGCGCATCCGCACATATTCT..... 102
682 alSerThrSerSerPheLeuGlyPheProProSerSerSerLeuLeu 698
101 TCGCCCAAGCAACGCGACT.....TC 82
699 ThrProAspSerAlaSerGlnrtrpGlntrp***ProAsnAlaLysGlySe 715
81 GGTAAATGGCGCGCGCTCATAAATGACTTGCCTCGCGTGTGCCCGGATGG 32
715 rValIleIleCysProAlaProSerSerAsnProGlyArgProAlaPheL 732
31 GCAGATT 24
732 euSerPhe 734

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seq_documentation_block:
ID AAW87504 standard; Protein; 1061 AA.
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XX AAW87504;
XX
XX 23-FEB-1999 (first entry)
DT
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914 oProGlySerThrAspSerThrSerAlaCysThrProThrProThrC 931
1018 GCGCGCGAGCGGCAACAATACTCC.....ATCAGCGG 1049
1031 yshiscysAlaGlyGlyLeu.SerValLeuThrPheHisProValThrAl 947
1050 CACCACCTCGGCGCATTCCTCAAAAACAACTCTTC..... 1086
1064 aThrAlaProGlySerProAlaProGlyGlyLeuTrpGlyThrAlaAlaG 964
1087 .....AAGTCACGACACCGCTCAACGGCGGCGGCGC 1119
1094 lyLeuTrpGlyTyrAlaGlnAlaThrValGlyAspTrpThrArg 980
1120 GCCATGTACCGATCGGCACTTATGCGCGTAAATGCGGTTGGACATCCT 1169
1181 SerAlaValProValGlyArgLysAlaSer..... 990
1170 GCCTACCTTGTCTTTCGCGGATTTAATCGTCGGGATFACCGACGCGCG 1219
1191 .....ArgaspProAlaProGlyAsp..... 997
1220 AGGCTTTGGTGTCTTGGAAATGGACGAAGACCTCGCTTTGTCAGC 1269
1298 .....Gly.SerProValTrpSerGlnLysCysGluLeuSerAlaTh 1011
1270 TTCGCTGCGCGGCAAAATACGAATACGCGCGCT.....GTTGCG 1310
1011 rGlnAla.ProSerGlnLeuAspSerLeuProAlaThrValArgValLys 1027
1311 CAAAGTGCTGGA 1322
1028 ArgGlnAlaGly 1031

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seq.name: /SIDSI/gcgdata/geneseq/geneseq-emb1/AA1999.DAT:AAW87503

seq_documentation_block:

XX AA87503 standard; Protein; 1212 AA.

XX AC AA87503;

XX DT 23-FEB-1999 (first entry)

XX DE Human N-methyl-D-aspartate receptor subunit encoded by clone NMDA22.

XX KW Human; N-methyl-D-aspartate receptor; NMDAR2C;

XX KW NMDA-activated cation-selective ion channel; glutamate receptor.

XX OS Homo sapiens.

XX PN US5849895-A.

XX PD 15-DEC-1998.

XX PF 20-APR-1994; 94US-02311193.

XX PR 20-APR-1994; 94US-02311193.

XX PR 20-APR-1993; 93US-0052449.

XX PA (SIBI-) SIBIA NEUROSCIENCES INC.

XX PI Daggett LP, Lu C;

XX DR WPI; 1999-069812/06.

XX DR N-PSDB; AAV82909.

XX PT DNA encoding N-methyl-D-aspartate receptor subunit - useful for the

XX PT assembly of functional glutamate receptor subunits

XX PS Example 3; Columns 253-262; 203pp; English.

XX PS The present sequence represents a human N-methyl-D-aspartate (NMDA)

CC

receptor subunit (NMDAR). The nucleic acid sequence does not contain the 366 5'-most nucleotides, by the insertion of 11 nucleotides between nucleotides 1300 and 1301, nor the 15 nucleotides at positions 1960-1974, nor the 1061 3' nucleotides, as set forth in AAV82889. The cDNA sequence is derived from clone NMDA21. The NMDAR subunits contribute to the formation of NMDA-activated cation-selective ion channels. In addition to being useful for the production of NMDA receptor subunit proteins, the nucleic acids are also useful as probes to identify and isolate nucleic acids encoding related receptor subunits. Functional glutamate receptors can be assembled from several NMDA receptor subunit proteins of one type (homomeric) or from combinations of subunit proteins of different types (heteromeric). The present invention also comprises methods for using such receptor subunits to identify and characterize compounds which affect the function of such receptors, e.g. agonists, antagonists and modulators of glutamate receptor function. The invention also comprises methods for determining whether unknown protein(s) are functional as NMDA receptor subunits.

XX Sequence 1212 AA;

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Ratio: 0.613 Gaps: 28

Percent Similarity: 35.306 Percent Identity: 23.265

alignment_block:

US-09-303-518D-131 x AA87503 ..

Align seg 1/1 to: AA87503 from: 1 to: 1212

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26 ATCTGCCATCGCGGACGAGCAAGTCATTATGACGGCGCGCC 75
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840 LeuAlaProSerArgIleGlyValAlaAlaValArgProHisArgPr 856

76 ATTACCGAAGTCGCGTGTCTGGCGAAGAATATGTCGGCATCGCCCTC 125
   : : : : : : : : : : : : : : : : : : : : : :
856' OProAlaArgProArgGlyLeuAlaPro...AlaHisAlaCysProPro. 871

126 GATGAAATCAAGGAGGTGAAGCGGTCAAAAAGG...CCAAGTGTGT 172
   : : : : : : : : : : : : : : : : : : : : : :
872 .....ProThrArgPro.GlnSerArgAlaProArgAlaG 883

173 TTGACAGACAAAAGATCGCGGTAGTATTACTCGCGCGCTTCAGCG 222
   : : : : : : : : : : : : : : : : : : : : : :
883 yAspArgGlnThrGlyValAlaArgLeu...CysAlaGlyLeuArg. 898

223 AAAATCGCGCTATTACCGTGGCGGAAAGCGCGT...ACTTCAGTCAGT 269
   : : : : : : : : : : : : : : : : : : : : : :
899 ..SerProArgAlaAlaProArgArgGlyArgProCysProThrSer 914

270 CGTGATTCGCGTGAAGCAACGCAAAATCGAGTTCGAACGCTACGTAC 319
   : : : : : : : : : : : : : : : : : : : : : :
915 ProGluCysArgAlaAlaGlnProGlyArgArgGlyArgCysGlyPr 931

320 CTGAAGCGCTGGCAAAATTTGAGCAGCGGAAAGAGTGGCGGCAACCTGATT 369
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931 oGlyThrAlaGlyGlyThrSerArgProSerGlyPro..... 944

370 CAATCAGGCTTATGACTCGCTTCGCCACCGCTCCGTTCCAGCAAAATCCC 419
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944 ..... 944

420 TGCCTAGATGCGGAGCGCGTTCGCCATCTTCGTCAATGCGATGGACACCA 469
   : : : : : : : : : : : : : : : : : : : : : :
944 ..... 944

470 ATCCGCTGGTGGCGGACCTACGTCATCATCAAGAACGCCCGCGAGAC 519
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945 .....CysArgPro.....ArgAlaV 950

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950 aThrThrAlaProPhe...LeuGluProThrAspProAla...AlaPro 964
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11779 ArgGlnAlaGly 1182
1311 CAAGTGTCTGGA 1322
seq_name: /SIDSL/gcgdata/geneseq/geneseq-emb1/AA2001.DAT:ABB60656
seq_documentation_block:
ID ABB60656 standard; Protein; 2088 AA.
XX AC ABB60656;
XX AC
XX 26-MAR-2002 (first entry)
XX DE Drosophila melanogaster polypeptide SEQ ID NO 8760.
XX KW Drosophila; developmental biology; cell signalling; insecticide;
XX KW pharmaceutical.
XX OS Drosophila melanogaster.
XX PN WO200171042-A2.
XX PD 27-SEP-2001.
XX PF 23-MAR-2001; 2001WO-US09231.
XX PR 23-MAR-2000; 2000US-191637P.
XX PR 11-JUL-2000; 2000US-0614150.
XX PA (PEKE) PE CORP NY.
XX PI Venter JC, Adams M, Li PWD, Myers EW;
XX WPI; 2001-656860/75.
XX DR N-PSDB; ABL04759.
XX PT New isolated nucleic acid detection reagent for detecting 1000 or more
XX PT genes from Drosophila and for elucidating cell signalling and cell-cell
XX PT interactions
XX PS Disclosure; SEQ ID NO 8760; 21pp + Sequence Listing; English.
XX CC The invention relates to an isolated nucleic acid detection reagent
XX CC capable of detecting 1000 or more genes from Drosophila. The invention is
XX CC useful in developmental biology and in elucidating cell signalling and
XX CC cell-cell interactions in higher eukaryotes for the development of
XX CC insecticides, therapeutics and pharmaceutical drugs. The invention
XX CC discloses genomic DNA sequences (ABL16176-ABL30511), expressed DNA
XX CC sequences (ABL01840-ABL16175) and the encoded proteins
XX CC (ABB57737-ABB72072).
XX CC The sequence data for this patent did not form part of the printed
XX CC specification, but was obtained in electronic format directly from WIPO
XX CC at ftp.wipo.int/pub/published_pct_sequences.
XX SQ Sequence 2088 AA;
alignment_scores:
Quality: 106.00 Length: 538
Ratio: 0.453 Gaps: 19
Percent Similarity: 43.494 Percent Identity: 20.260
alignment_block:
US-09-303-518D-131/rev x ABB60656
Align seg 1/1 to: ABB60656 from: 1 to: 2088
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1293 TTCGTATTTGCCGGGACAGCAAGCTGCACAAAGCGAGGTCTTCTCGT 1244

950 aThrThrAlaProPhe...LeuGluProThrAspProAla...AlaPro 964
570 TGTGTGTAAGACGACGCGCAGCTGCGCTGAAATGC...TGCCA 616
11779 ArgGlnAlaGly 1182
1311 CAAGTGTCTGGA 1322
seq_name: /SIDSL/gcgdata/geneseq/geneseq-emb1/AA2001.DAT:ABB60656
seq_documentation_block:
ID ABB60656 standard; Protein; 2088 AA.
XX AC ABB60656;
XX AC
XX 26-MAR-2002 (first entry)
XX DE Drosophila melanogaster polypeptide SEQ ID NO 8760.
XX KW Drosophila; developmental biology; cell signalling; insecticide;
XX KW pharmaceutical.
XX OS Drosophila melanogaster.
XX PN WO200171042-A2.
XX PD 27-SEP-2001.
XX PF 23-MAR-2001; 2001WO-US09231.
XX PR 23-MAR-2000; 2000US-191637P.
XX PR 11-JUL-2000; 2000US-0614150.
XX PA (PEKE) PE CORP NY.
XX PI Venter JC, Adams M, Li PWD, Myers EW;
XX WPI; 2001-656860/75.
XX DR N-PSDB; ABL04759.
XX PT New isolated nucleic acid detection reagent for detecting 1000 or more
XX PT genes from Drosophila and for elucidating cell signalling and cell-cell
XX PT interactions
XX PS Disclosure; SEQ ID NO 8760; 21pp + Sequence Listing; English.
XX CC The invention relates to an isolated nucleic acid detection reagent
XX CC capable of detecting 1000 or more genes from Drosophila. The invention is
XX CC useful in developmental biology and in elucidating cell signalling and
XX CC cell-cell interactions in higher eukaryotes for the development of
XX CC insecticides, therapeutics and pharmaceutical drugs. The invention
XX CC discloses genomic DNA sequences (ABL16176-ABL30511), expressed DNA
XX CC sequences (ABL01840-ABL16175) and the encoded proteins
XX CC (ABB57737-ABB72072).
XX CC The sequence data for this patent did not form part of the printed
XX CC specification, but was obtained in electronic format directly from WIPO
XX CC at ftp.wipo.int/pub/published_pct_sequences.
XX SQ Sequence 2088 AA;
alignment_scores:
Quality: 106.00 Length: 538
Ratio: 0.453 Gaps: 19
Percent Similarity: 43.494 Percent Identity: 20.260
alignment_block:
US-09-303-518D-131/rev x ABB60656
Align seg 1/1 to: ABB60656 from: 1 to: 2088
1343 CAGCCTTCCTTCTCAATGTTTCCAGCACTTTGCGCAACAGCGGCCGTA 1294
1297 LysProSerValGluThrThrThrProAlaProThrSerAlaGlnPh 1313
1293 TTCGTATTTGCCGGGACAGCAAGCTGCACAAAGCGAGGTCTTCTCGT 1244

1313 eserPheGlyPheGlyGlnSerAsnGlnGlyLysAspValAlaAspSerL 1330
1243 CCAATTCACCAACCAACCAACCTGCGGCTGTGCTGATCGCCGACGATT 1194
1330 ysLysThrGluAlaProLysThrPheMetPheGlyValSer..... 1343
1193 AAATCGCCCAACCAAGTAGGAGGATGTCACACGGCATTCGCGCTC 1144
1344 LysValGluGluProLysThrValSerPheGlyThrGlyIleLysGluTh 1360
1143 ATAAGTCGCGATCGATACCATCGCGCTGCGCGCTGACGGCTGTCG 1094
1360 rThrAlaThrSerSerThrGluAlaThrAlaProThrProAlaAlaAla 1377
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1377 laProAlaProValGlnPheValPheLysAlaProThrThrAlaThrThr 1393
1043 ATGGAGTATTGTC..... 1029
1394 AlaSerSerLeuThrThrThrIleSerThrThrSerAsnAlaProAlaLe 1410
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1410 uGlyGlyPheSerPheGlyAlaProSerSerSerThrValSerSers 1427
982 CGATAACGGAATCGATTGTTGGTAGCTCCCAATATCATCGCGCT 933
1427 erThrThrSerThr.....SerAlaAsnPro 1435
932 TGTGCAATCGCACCGTTC..... 915
1436 AlaAlaValLysProMetPheSerTrpSerGlyAlaGlySerAlaValSe 1452
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1452 rSerThrSerSerGlnGlnProValAlaLysAlaProThrLeuGlyP 1469
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1469 heGlyValSerSerThrValThrThrThrThrThrThrThrThrThr 1485
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1486 PheAlaPheThrProAlaSerGlyLeuAspProAlaAlaAlaThrSerAl 1502
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1502 aProAlaAlaGlyAlaGlyPheSerPheGlySerGlnSerLysProAlaT 1519
835 CGGTACGC.....AAGAGCGCGGTTTGTG 810
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809 ACTTCAGCGCCCAACCAACCAACCGCTGCTGATTCAGACGCGCTGT 760
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1622 AlaLysSerAsnSerAlaAla.....ValGlySerAlaAla..... 1633
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374 GATTGAATCAGGTTGCGGCGCACTTTTTCGCTGCTCAATTTTGGCAGCGC 325
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177 TTCAAACAGC.....ACTGGCCCTTTT 155
1697 lGlyAsnSerLeuAlaGlyValGlyAlaProValAlaAlaThrThrProAla 1714
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1731 AlaAlaAlaAlaProValPheGlySerGlySerThrIleProSerAlaGl 1747
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1747 yPheGlyAlaProAlaAlaAlaAlaProLeuAlaAlaProAlaLeuProG 1764
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seq_name: /SIDS1/gcgdata/geneseq/geneseq-emb1/AA2001.DAT:AAU48589
seq_documentation_block:
ID AAU48589 standard; Protein; 372 AA.
XX
AC AAU48589;
XX
DT 27-FEB-2002 (first entry)
XX
DE Propionibacterium acnes immunogenic protein #9485.
XX
KW SAPHO syndrome; synovitis; acne; pustulosis; hypervellitis;
KW uveitis; endophthalmitis; bone; joint; central nervous system; ELISA;
KW inflammatory lesion; acne vulgaris; enzyme linked immunosorbent assay;
KW dermatological; osteopathic; neuroprotectant.
XX
OS Propionibacterium acnes.
XX
PN WO200181581-A2.


```

322 CAGGTACGTAGCTGCGTTCGAACGTGATTGGTGGTCCCTTCA...ACGGCA 276
||||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
239 leGlyThrThrArgArgGly.....SerProileProSerLeuThrala.253
||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
275 ATCAGCAGTACTGAAGTACGGCGTTCGCGCACGGTGAATA..... 234
||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
254 SerThrIleValArgSerArgSerValArgTrpValArgSerSe 270
||||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
233 ...CGCGGATTTTGCCTGAAGCGCGCAGTAAATACCTACGCCGGAT 188
||||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
270 rSerSerProHisProArgThrSerProThrGlyThrIleGlys 287
||||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
187 TC.....TTTGTCTTCAACAGCAGTTCGCCCTTTTTCGACGGCTTCA 144
||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
287 erThrArgPheArgProSerAlaArgSerTrp..... 297
||||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
143 CCTTCCTTGATTTCATCGAGGGCGCATGCCGACATATTTTCGCCCAAG 94
||||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
298 .....SerSerProse 301
93 CAACGCGACTTCG.....GTAATGCGCGG 69
||||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
301 rSerAlaThrThrSerArgAsnGlyValMetThrGly 313
||||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||

```

seq_name: /SIDS1/gcgdata/geneseq/geneseq-emb1/AA11994.DAT:AA44890

seq_documentation_block:

ID AAR44890 standard; Protein; 532 AA.

XX AC AAR44890;

XX DT 22-JUN-1994 (first entry)

XX DE Diphtheria toxin delta-142-147-148 mutant.

XX KW DT; protein exotoxin; NAD-dependent ADP-ribosyltransferase; vaccine;
 KW diphtheria toxin; deletion mutant; mutagen; variant; double mutant;
 KW reversion mutation; site-directed mutagenesis.

XX OS Corynebacterium diphtheriae.

XX FH Key Location/Qualifiers

FT Protein

FT 1..532

FT /note= "Diphtheria toxin mutant; Glu(142), Val(147)
 and Glu(148) have been deleted"

FT WO9325210-A.

PN 23-DEC-1993.

XX PF 17-MAY-1993; 93WO-US04606.

XX PR 18-JUN-1992; 92US-0901712.

XX PA (HARD) HARVARD COLLEGE.

XX PI Collier RJ, Killeen K, Mekalanos J;

XX WPI; 1994-007178/01.

XX N-PSDB; AAQ54338.

XX New DNA encoding diphtheria toxin deletion mutants - with no
 toxicity and low risk of reversion, and derived toxoids and
 transformed cells, useful in vaccines

XX Claim 11; 42pp; English.

XX Oligonucleotide-directed mutagenesis of the wild-type diphtheria
 gene (specifically the region encoding the DT-A fragment) results
 CC in deletion of the codons for Val-147 and active site residue
 CC Glu-148. The resulting mutagen is not toxic, making it useful in
 CC diphtheria vaccines. The risk of reversion to toxicity is much

CC lower for the 147-148 double mutant than for the prior art 148
 CC single mutant, while its immunogenicity is not impaired. The
 CC 147-148 mutagen opt. has other amino acid residues subst. or
 CC deleted, e.g. wild-type Glu(142) is deleted. The specification
 CC includes the wild-type DT amino acid sequence (see AAR44888) but does
 CC not include any mutant sequences; the wild-type sequence was modified
 CC according to the description in the claims to give AAR44890.
 XX
 SQ Sequence 532 AA;

alignment_scores:

Quality: 105.00 Length: 461
 Ratio: 0.536 Gaps: 25
 Percent Similarity: 42.516 Percent Identity: 20.390

alignment_block:

US-09-303-518D-131 x AAR44890 ..

Align seg 1/1 to: AAR44890 from: 1 to: 532

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31 CCCATCGCGGCGAGACCGGAGCAAGTCATT...TATGACGCGCGCGCCAT 77
||||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
72 ProLeuSerGlyLysAlaGlyGlyValLysValThrTyrProGlyLe 88
||||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
78 TACCGAAGTCGCGTCTGCTTGGCGCAAGAATATGTCGCATGCGCCCTCGA 127
||||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
88 uThrLysValLeu.....AlaL 94
||||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
128 TGAATAATCAAGGAAGGTGAAGCGGTCAAAAAA..... 159
||||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
94 eulysValaspasnAlaGluThrIleLysLysGluLeuGlyLeuSerLeu 110
||||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
160 .....GCCCAAGTCTGCTTTTGAAGACAAAAA 185
||||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
111 ThrGluProLeuMetGluGlnValGlyThrGluGluPheIleLysArgPh 127
||||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
186 GAATCGCGG.....GTAGTATTACTGCGCGGCTTCAGGCAAAA 226
||||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
127 eGlyAspGlyAlaSerArgValValLeuSerLeuProPheAlaGlySerS 144
||||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
227 TCGCGCTATTCAACGTCGCGGCAAGCGGTACTTTCAGTCAGTCGTCGATT 276
||||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
144 eSerTyrIleAsnAsnTrpGluGlnAlaLysAlaLeuSerValGluLeu 160
||||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
277 GCGGTGAAGGCAACGACCAATCGAGTTCGAA..... 309
||||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
161 .....GluIleAsnPheGluThrArgGlyLysArgGl 171
||||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
310 .....CGCTACGTACCTGAAGCGCTGGCAAAATGAGCA 343
||||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
171 yGlnAspAlaMetTyrGluTyrMetAlaGlnAlaCysAla.....G 185
||||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
344 GCGAAAGTGCAGCGCAACCTGATTCATCAGGCTTA..... 381
||||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
185 lYasnArgValArgArgSer...ValGlySerSerLeuSerCysIleAsn 200
||||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
382 .....TGGACTGCGCTTCGCGACCGCTCCGTTTCAGCAAAATCCCTGCCGT 425
||||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
201 LeuAspTrpAspValIleArgAspLysThrLysThrLysIleGluSerLe 217
||||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
426 AGATGCGGAGCGGTTTCGCTCAATGCGATGGACACCAATCCGC 475
||||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
217 u.....LysGluHisGlyProIleLysAsnLysMetSerGluSerProA 232
||||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
476 TGGCTGCCGACCCCTACGGTCATCATCAAGAAGCGCGCAAGACTTCAA 525
||||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
232 snLysThrValSerGluGluLysAlaLysGlnTyrLeuGluGluPheHis 248
||||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
526 CGCGGCTGTTGGTATTGAGCGCGCTGACCGACGCTAAATCCATGTGTG 575
||||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
249 GlnThrAlaLeuGluHisProIleuSerGluLeuLysThr..... 262
||||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||

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576 TAAAGCAGCAGCGCGCAGACGTCGCGTCTGAAATGCTGCAATATCGAAA 625
262 .....
626 CACATGAATTGGCGCGCCGATCCTCCGGCTTGAGTGGCAGCAGCATT 675
263 .....
676 CATTTTCATGAGCGAGTGGCGGGAATAAAGCGGTGGACCAATCAATTA 725
269 ValPhe.....AlaGlyAlaAsnTyrAlaAlaTyrAlaValAsnVa 282
726 TCAAGACGTGATT..... 738
282 lAlaGlnValIleAspSerGluThrAlaAspAsnLeuGluLysThrThra 299
739 .....GCTATCGGAGCTTGG.....TTCGTAACA 762
299 laAlaLeuSerIleLeuProGlyIleGlySerValMetGlyIleAlaAsp 315
763 GGCGGTCTG.....AATACCGAGCGGTGGTGGCCCTGGCGCGCTGCA 806
316 GlyAlaValHisAsnThrGluGluIleValAla..... 327
807 AGTCAACAACCGCGCTCTTGGTACCGTGGTGGTGGCGAAGTGTC 856
328 .....GlnSerIleAlaLeuSerSerLeuMetValAlaGlnAlaIleP 342
857 AACTTACCGCGCGGAAATGGTTCACCGGACACCGCGGTGATTCGGT 906
342 roLeu...ValGluLeuValAsp..... 349
907 TCGGTATTGAACGGTGGCGATTCACAGCGCGCGATGATTATTGGAGC 956
350 .....IleGlyPheAlaAlaTyrAsnPheValGluSe 360
957 CTACCAACATCAGATTCCGTTATTCGAAGAGCGCGCAGCAAGAGCTGT 1006
360 rIleIleAsnLeuPheGlnValHisAsnSerTyrAsnArgProAlaT 377
1007 TCGGCTGGTGGCGCG.....CAGCG.....GACAAA 1035
377 yr.....SerProGlyHisLysThrGlnPropheLeuHisAspGly 390
1036 TACTCCATFACGCGCAGCAGCTCTC..... 1059
391 TyrAlaValSerTyrAsnThrValGluAspSerIleIleArgThrGlyPh 407
1060 .....GGCCATTCTCTAAACAACTCTTCAAGTTCACGA 1096
407 eGlnGlyGluSerGlyHisAspIleLys.....IleT 418
1097 CAGCGCTCAACGGCGCGCGCGCATGGTACCGATCGGCAGCTATGAG 1146
418 hrAlaGluAsn.....ThrProLeuProIleAlaGlyVal... 429
1147 GCGGTAATGCGGTGGATGATCCTGCTACCTG 1179
430 .....LeuLeuProThrIle 434
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seq_name: /STDS1/gcdata/geneseq/geneseq-emb1/AA2001.DAT:ABG21970

seq_documentation_block:

ID ABG21970 standard; Protein; 575 AA.

XX AC ABG21970;

XX DT 18-FEB-2002 (first entry)

XX DE Novel human diagnostic protein #21961.

XX DE Human; chromosome mapping; gene mapping; gene therapy; forensic;

KW food supplement; medical imaging; diagnostic; genetic disorder.

```
XX Homo sapiens.
XX WO200175067-A2.
XX 11-OCT-2001.
XX 30-MAR-2001; 2001WO-US08631.
XX 31-MAR-2000; 2000US-0540217.
XX 23-AUG-2000; 2000US-0649167.
XX (HYSE-) HYSEQ INC.
XX Drmanac RT, Liu C, Tang YT;
XX WPI; 2001-639362/73.
XX N-PSDB; AAS86157.
XX New isolated polynucleotide and encoded polypeptides, useful in
XX diagnostics, forensics, gene mapping, identification of mutations
XX responsible for genetic disorders or other traits and to assess
XX biodiversity.
XX Claim 20; SEQ ID NO 52329; 103pp; English.
XX The invention relates to isolated polynucleotide (I) and
XX polypeptide (II) sequences. (I) is useful as hybridisation probes,
XX polymerase chain reaction (PCR) primers, oligomers, and for chromosome
XX and gene mapping, and in recombinant production of (II). The
XX polynucleotides are also used in diagnostics as expressed sequence tags
XX for identifying expressed genes. (I) is useful in gene therapy techniques
XX to restore normal activity of (II) or to treat disease states involving
XX (II). (II) is useful for generating antibodies against it, detecting or
XX quantitating a polypeptide in tissue, as molecular weight markers and as
XX a food supplement. (II) and its binding partners are useful in medical
XX imaging of sites expressing (II). (I) and (II) are useful for treating
XX disorders involving aberrant protein expression or biological activity.
XX The polypeptide and polynucleotide sequences have applications in
XX diagnostics, forensics, gene mapping, identification of mutations
XX responsible for genetic disorders or other traits to assess biodiversity
XX and to produce other types of data and products dependent on DNA and
XX amino acid sequences. ABG00010-ABG30377 represent novel human
XX diagnostic amino acid sequences of the invention.
XX Note: The sequence data for this patent did not appear in the printed
XX specification, but was obtained in electronic format directly from WIPO
XX at ftp.wipo.int/pub/published_pct_sequences.
XX SQ Sequence 575 AA;
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alignment_scores:

Quality: 105.00 Length: 519
Ratio: 0.515 Gaps: 24
Percent Similarity: 39.306 Percent Identity: 19.653

alignment_block:

US-09-303-518D-131 x ABG21970 ..

Align seq 1/1 to: ABG21970 from: 1 to: 575

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31 CCCATCGCGGGCAGACCGGAGCAAGTCATTATGAGCGCCGCGCATTCAT 80
   ||| :|||:|:| ||| ||| ||| ||| ||| ||| ||| ||| |||
46 ProGlySerGlyLysSerProProValAlaSerGlyGlyProAla..... 60
81 CGAAGTCGCGCTTCTGTTGGCGAAGAAATATGTCGCGCATGCGCCCTCGATGA 130
   :|||
61 .....ArgAlaLysProHisArgA 67
131 AAATCAAGGAAGGTGAAGCGCGTCAAAAAGGCCAAGCTGCTGTTTGAAGAC 180
   :|:| :|:| :|:| :|:| :|:| :|:| :|:| :|:| :|:| :|:|
67 laValHisValSerProAlaProSerGlyGly..... 77
```



```

877 777 ..... 877
716 CCATCAATTATCAAGACGTGATTGCTATCGAGCTTTGTTCTGTAACAGC 765
878 .....ProArgp 880
766 GGTCTGAATACCGAGCGCGGTGGTTCCTTGGCGGCTGCAAGTCAACAA 815
880 roSerLeuGlyProAlaArgCysProLeuGlyAlaProAlaProAla 896
816 ACCGCGCTCTTGGCTACCGTTCCTTGGTGGGAAGTGTCTCAACTTACG 865
897 ProAlaProThrAlaThrArg .....ProAlaGlyAlaThrPAr 909
866 CCGCGCAATTGGTTGACGCGGCAACCGCGTGATTTCGCGTTTCGTTATG 915
909 gArgArg .....S 912
916 AACGGTGGGATTCACAAAGCGCGCATGATTATTTGGGACGCTACCAAA 965
912 erArgCysAlaCys ..... 916
966 TCAGATTTCGTTATCGAAGCGCGCAGCAAGAGCTGTTTCGCTGGG 1015
917 .....ArgSerThrGlyArgProAlaArgAlaSerArgGlnG 930
1016 T ..... 1016
930 yProProGlySerThrAspSerThrSerAlaCysThrProThrProt 947
1017 ..TGCGCGCGAGCGGACAAATACTCC .....ATCAC 1046
947 hrCysHisCysAlaGlyGlyLeu.SerValLeuThrPheHisProValTh 963
1047 GCGCACCACTCTCGGCCATTTCCTAAACCAACACTCTTCAAGTTCACGA 1096
963 rAlaThrAlaProGlySerProAlaProGlyGlyLeuThrGlyThrAlaA 980
1097 CAGCGGTACAGGGGGACCGGCCATGGTACCGATCGGCACCTTATGAG 1146
980 laGlyLeuThrGlyTrpAlaGlnAlaThrGluThrValGlyAspThrP 996
1147 CGC .....GTAATCGCTTGGACATCTCGCTACCTTTCCTTTCGCGCA 1190
997 ArgSerAlaVal**ProValGlyArgLysAlaSer .....ArgAs 1010
1191 TTTAATCGTCGCGCATACCGACAGCGCGAGCTTTGGTTGCTTGGAA 1240
1010 pProAlaProGlyAsp .....Gly.SerProVal 1019
1241 TGGACGAAGAAGACCTCGCTTTGTCAGCTGCTGCGCGCGGCAATATAC 1290
1020 TrpSerGlnLysCysGluLeuSerAlaThrGlnAla.ProSerGlnLeuA 1036
1291 GAATACGGCCGCT .....GTTCGCAAGTGTGGA 1322
1036 spSerLeuProAlaThrValArgValLysArgGlnAlaGly 1049

seq_name: /STDS1/gcgdata/geneseq/geneseq-emb1/AA2000.DAT:AA26239
seq_documentation_block:
ID AAB26239 standard; Protein; 1232 AA.
XX
AC AAB26239;
XX
XX
DT (first entry)
XX
DE Human N-methyl-D-aspartate receptor subunit NMDAR1A #17.
XX
KW Human; N-methyl-D-aspartate receptor; NMDA; NMDAR1A; ionotropic;
KW glutamate receptor; drug screening; animal model; disease diagnosis;
KW genetic screening.

```

PI Daggett LP, Lu C;
 XX WPI; 2000-578607/54.
 DR N-PSDB; AAA95032.
 XX
 PT Novel DNA fragment encoding human N-methyl-D-aspartate receptor subunit
 PT for identifying mutations and for developing drugs against various
 PT disease states
 XX
 XX Example 3; column 255-264; 205pp; English.
 XX
 CC The present sequence is a subunit (designated NMDAR1A) of the human
 CC N-methyl-D-aspartate (NMDA) receptor. This is an ionotropic glutamate
 CC receptor which contains cation-specific ligand-gated ion channels. The
 CC protein and its coding sequence can be used in disease diagnosis and in
 CC research to identify other, similar proteins. They can also be used as
 CC probes, for example in genetic screening, and in drug screening, as well
 CC as enabling the production of animal disease models.
 XX
 XX Sequence 1232 AA;

alignment_scores:

Quality: 105.00 Length: 483
 Ratio: 0.593 Gaps: 28
 Percent Similarity: 36.646 Percent Identity: 23.602

alignment_block:

US-09-303-518D-131 x AAB26239 ..

Align seg 1/1 to: AAB26239 from: 1 to: 1232

26 ATCTGCCATCGCGGACAGCGGACGCAAGTCATTATGACGGCCGCGCC 75
 856 LeuAlaProSerArgIleGlyValAlaAlaValArgProHisArgPR 872
 76 ATTACCGAAGTCGGCTTCTGGCGAAGAATATGCGGCATCGCCCGCTC 125
 872 oProAlaArgProArgGlyLeuAlaPro...AlaHisAlaCysProPro. 887
 126 GATGAATCAAGGAGTGAAGCGGTCAAAAAGG...CCAAGTGGTGT 172
 888ProThrArgPro.GlnSerArgAlaProArgAlaGly'899
 173 TTGAAGACAAAGAATCGCGGTAGTATTACTGCGCGCTTCAGGC 222
 899 yAspArgGlnThrGlyValAlaArgLeu...CysAlaGlyLeuArg. 914
 223 AAAATCGCGCTATTTCACGTGGCGGAAAGCGCGT...ACTTCAGTCAGT 269
 915 ..SerProArgAlaAlaProArgArgGlyArgProCysProThrSer 930
 270 CGTGATTCGGTGAAGCAGCAGCAAGTTCGAGTTCGAACGCTACGTAC 319
 931 ProGluCysArgAlaAlaGlnProGlyArgArgGlyGlyArgCysGlyPr 947
 320 CTGAAGCGCTGGCAAAATAGCAGCAGAAAGTTCGCGCGCAACCTGATT 369
 947 oGlyThrAlaGlyGlyThrSerArgProProSerGlyPro..... 960
 370 CAATCAGGCTTATGACTCGCGTTCGCACCCCTCGCTTCACCAAAATCCC 419
 960
 420 TGGCGTAGATCGGAGCCGCTTCGCCATCTCTGTCATGCGATGGACACCA 469
 960
 470 ATCCGCTGGCTCCGACCCCTACCGTCAATCAAAAGAGCCGCGCAAGAC 519
 961CysArgPro.....ArgAlaV 966
 520 TTCAACAGCGCGCTGTGTGTTATGTAGCCGCTGACCGACGTAATAATCCA 569

966 alThrAlaProPhe...LeuGluProThrAspProAla...AlaPro 980
 570 TGTGTGTAAGCAGCAGCGCAGCGTCCGCTCTGAAAATGC...TGCCA 616
 981 SerSerArgSerSerArgSerProArgSerTrpArgThrCysArgCysSe 997
 617 ATATCGAAACACATGAATTTGG.....CGGCCCGCATCC..... 650
 997 rValArgSerSer.....TrpProGlyGlyArgProCys***ThrArgP 1012
 651TGCCGGCTTGAGTGG 665
 1012 roGlyProGlyAlaArgAlaArgValThrLeuProCysProAlaProTrp 1028
 666 CAGGCACATTCATTCATCGACCGCAGTCGCGCGAATAAAACCGGTGGA 715
 1028 1028
 716 CCATCAATTATCAAGACGTGATTGTCGTATCGGACGTTTGTCTGAACAGGC 765
 1029ProArgP 1031
 766 CGTCTGAATACGAGCGCGCTGTGCTTGGCGCGCTGCCAAGTCAACAA 815
 1031 roSerLeuGlyProAlaArgCysProLeuGlyAlaProAlaProAla 1047
 816 ACCGCGCTCTTGGTACCGTTCGCGGAGGTTCTCAACTACCG 865
 1048 ProAlaProThrAlaThrArg.....ProAlaGlyAlaTrpAr 1060
 866 CCGCGCAATTTGTTGACGCGGACAAACCGCGTATTTCGGTTCGGTATTG 915
 1060 gA'gArg.....S 1063
 916 AAGCGTTCGATTTCACAAAGCGCGCATGATTATTTGGACGCTACCACAA 965
 1063 erArgCysAlaCys..... 1067
 966 TCAGATTTCGTTATCGAAGAGCGCGCAAGAGCTGTTTCGGCTGGG 1015
 1068ArgSerThrGlyArgProAlaArgAlaSerArgGlnG 1081
 1016 T..... 1016
 1081 yProProGlySerThrAspSerThrSerAlaCysThrProThrProT 1098
 1017 ..TGCGCGCAGCGCGCAAAATACTCC.....ATCAC 1046
 1098 hrCysHisCysAlaGlyGlyLeu.SerValLeuThrPheHisProValTh 1114
 1047 GCGCACCACTCTCGGCCATTTCTCTAAAAACAACTCTTCAAGTTCACGA 1096
 1114 rAlaThrAlaProGlySerProAlaProGlyGlyLeuTrpGlyThrAlaA 1131
 1097 CAGCGCTCAACGCGCGCGCATGTCACCGATCGGCACCTTATGAG 1146
 1131 laGlyLeuTrpGlyTrpAlaGlnAlaThrGluThrValGlyAspTrpThr 1147
 1147 CGC.....GTAATGCGTTGGACATCTCTGCCTACCTTGTCTTTCGCGCA 1190
 1148 ArgSerAlaVal**ProValGlyArgLysAlaSer.....ArgAs 1161
 1191 TTTAATCGTCGGCATACCGACAGCGCGGAGGCTTTGGTGTGTTGGAAT 1240
 1161 pProAlaProGlyAsp.....Gly.SerProVal 1170
 1241 TGGACGAAGAGACCTCGCTTGTGACAGCTTCGTCTGCCCGGCAATAC 1290
 1171 TrpSerGlnLysCysGluLeuSerAlaThrGlnAla.ProSerGlnLeuA 1187
 1291 GAATACGCGCGCT.....GTTGCGCAAGTGTGGA 1322

Mon Jul 1 09:25:33 2002

us-09-303-518d-131.rag

Page 57

1187 spSerLeuProAlaThrValArgVallysArgGlnAlaGly 1200

